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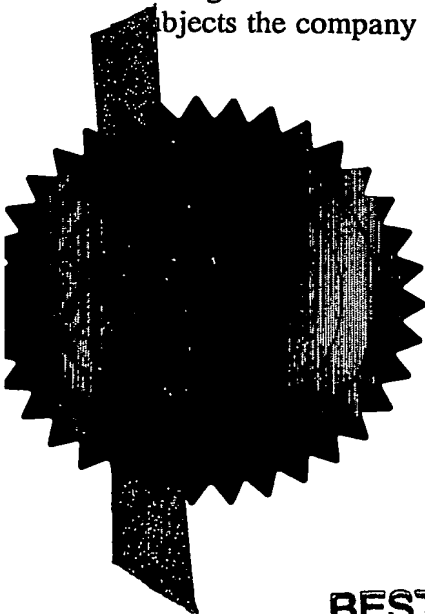
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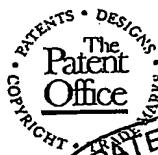
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The Patent Office

Cardiff Road
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NP10 8QQ

1. Your reference

P015069GB AAW

2. Patent application number

(The

- 6 SEP 2002

3. Full name, 0220809.8
each applicant (underline all surnames)

University College London
Gower Street
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United Kingdom

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

59 506 39 004 A
798652002

4. Title of the invention

Immune Modulator

5. Name of your agent (if you have one)

D Young & Co

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

21 New Fetter Lane
London
EC4A 1DA

Patents ADP number (if you know it)

59006

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
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Date of filing
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Number of earlier application

Date of filing
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8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

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Description 68

Claim(s) 4

Abstract 1

Drawing(s) 10 + 10 

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Statement of inventorship and right to grant of a patent (Patents Form 7/77) 0

Request for preliminary examination and search (Patents Form 9/77) 0

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Any other documents (please specify) None

11. I/We request the grant of a patent on the basis of this application.
Signature D. Young & Co Date 06 September 2002
D Young & Co (Agents for the Applicants)

12. Name and daytime telephone number of person to contact in the United Kingdom James P Tanner 023 8071 9500

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Immune modulator

FIELD OF THE INVENTION

- 5 The present invention relates to immune modulators, in particular vaccines that modulate a cellular immune response and uses thereof.

BACKGROUND TO THE INVENTION

- 10 Vaccines and other immune modulators have a major impact in reducing morbidity and mortality from disease. The primary immunity elicited by most current vaccines appears to be mediated by the humoral immune response. For diseases that may require a cellular immune response, such as tuberculosis and leishmaniasis, there are currently no available vaccines that are uniformly effective.

- 15 Typically, adjuvants are added to vaccines. The role of the adjuvant is to enhance the body's immune response to specific antigens of the vaccine. Commonly used adjuvants typically produce a humoral immune response but not a cell-mediated immune response. In addition, aluminium adjuvants for example may cause negative
20 side effects, such as sterile abscesses, erythema, swelling, subcutaneous nodules, granulomatous inflammation and contact hypersensitivity.

- A vaccine or other immune modulator is sought that modifies a cellular immune response and in particular the T helper cell response, for example, the T helper cell 1
25 (Th1) and T helper cell 2 (Th2) response.

SUMMARY OF THE INVENTION

- 30 From birth to death the immune system is educated, constantly stimulated and regulated through contact with the environment. Modern urbanisation and public health measures to prevent infectious disease have virtually eliminated this exposure

leading to an unprecedented rise in diseases – such as allergies and neoplastic diseases. Restoring the beneficial affects of the environment through the use of killed suspensions of harmless beneficial environmental bacteria may redress the normal balance of the immune system thus acting, therapeutically and/or prophylactically in the treatment of diseases and/or in promoting a healthy immune system.

Accordingly, the present invention is predicated upon the surprising finding that a whole cell of a bacterium from any one of the actinomycete genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* administered to a test subject can elicit a modification of the immune system, in particular the cellular immune system of that test subject.

The phrase "cellular immune system", as used herein, includes a cell-mediated immune response which depends upon the presence of T lymphocytes. The term "T lymphocytes" includes cytotoxic T lymphocytes, helper T cells, suppresser T cells and regulatory T cells. Modification of a cell-mediated immune response may be used, for example, to overcome cell-mediated immune disorders including, for example an immune system imbalance and immune hypersensitivity.

The terms "modulate", "modify", "modification" and other derivatives thereof, as used herein, mean downregulating, inhibiting, inducing, stimulating, upregulating, altering or otherwise affecting a component or components of the cellular immune system.

DETAILED ASPECTS OF THE PRESENT INVENTION

In one aspect, the present invention provides an immune modulator composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*.

The term "immune modulator", as used herein, means a substance which modulates a cellular immune system of a subject.

The term "whole cell", as used herein, means a bacterium which is intact, or substantially intact. In particular, the term "intact" as used herein means a bacterium which is comprised of all of the components present in a whole cell, particularly a whole, viable cell, and/or a bacterium which has not been specifically treated to remove one or more components from it. By the term "substantially intact" as used herein it is meant that although the isolation and/or purification process used in obtaining the bacterium may result in, for example, a slight modification to the cell and/or in the removal of one or more of the components of the cell, the degree to which such a modification and/or removal occurs is insignificant. In particular, a substantially intact cell according to the present invention has not been specifically treated to remove one or more components from it.

Although it has been suggested that individual components of bacterial cells could be used to elicit an adjuvant effect, prior to the present invention the use of whole cells of bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* to modulate a cellular immune response was not contemplated. Surprisingly, it has been found that by using a whole cell of a bacterium from said genera, modulation of a cellular immune system can be effected. The modulation of a cellular immune response caused by administration of said whole cell of said bacterium may be advantageously long lasting as compared with the response elicited by administration of an individual component of the bacterium.

Preferably, the composition according to the present invention comprises more than one whole cell, and more preferably comprises a plurality of whole cells.

In a further aspect, the present invention provides an immune modulator composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, which immune modulator composition in use modifies a cellular immune response.

In another aspect, the present invention provides an immune modulator composition comprising an antigen and an adjuvant, wherein said adjuvant comprises a whole cell

of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*.

5 In another aspect, the present invention provides a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* and optionally a pharmaceutically acceptable carrier, diluent or excipient, which immune modulator composition in use modifies a cellular immune response.

10 The present invention yet further provides a process of preparation of a pharmaceutical composition of the present invention, said process comprising admixing one or more of the compounds of the present invention with a pharmaceutically acceptable diluent, excipient or carrier.

15 In a further aspect, the present invention provides an immune modulator composition and/or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, and at least one antigen or antigenic determinant.

20 Suitably, the antigen or antigenic determinant may be an antigen or antigenic determinant from one or more of the following: BCG (bacillus of Calmette and Guerin) vaccine, diphtheria toxoid vaccine, diphtheria/tetanus/pertussis (DTP or Triple) vaccine, pertussis vaccine, tetanus toxoid vaccine, measles vaccine, mumps vaccine, rubella vaccine, OPV (oral poliomyelitis vaccine) and *Mycobacterium*
25 *vaccae*, or part thereof (as taught in GB0025694.1).

Suitably, the immune modulator composition and/or pharmaceutical composition may comprise two or more such antigens or antigenic determinants.

30 In another aspect, the present invention provides an immune modulator composition comprising an antigen or an antigenic determinant and an adjuvant, wherein said

antigen or antigenic determinant comprises a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*.

When it is the case that whole cell of the bacterium functions as an antigen or antigenic determinant the composition may suitably comprise at least one, preferably at least two, more preferably at least three, further antigens or antigenic determinants.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of one or more of an infection (eg. a bacterial, viral, or parasitic infection, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, for example, major trauma stress, psychosocial stress and chronic stress; an allergy (eg. asthma including allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses)), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (eg. an immune system imbalance in children and the elderly). An immune system imbalance in the elderly may be referred to as immunosenescence.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of one or more of a parasitic infection, such as, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis.

30

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera

Rhodococcus, Gordonia, Nocardia, Dietzia, Tsukamurella and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of stress, such as, for example, major trauma stress, psychosocial stress and/or chronic stress.

- 5 The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus, Gordonia, Nocardia, Dietzia, Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of one or more of asthma (including allergic asthma), hayfever, allergic dermatitis (eczema),
10 anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges for instance *Culicoides* (which causes Sweet Itch in horses). Preferably, the immune modulator composition or pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus, Gordonia, Nocardia, Dietzia, Tsukamurella* and *Nocardioides*, is used
15 in the manufacture of a medicament for the treatment or prevention of asthma including allergic asthma, and allergies to insect bites – such as midges for instance *Culicoides* (which causes Sweet Itch in horses).

- The present invention further provides the use of an immune modulator composition or
20 a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus, Gordonia, Nocardia, Dietzia, Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of one or more of heaves and/or COPD, particularly in horses.

- 25 The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus, Gordonia, Nocardia, Dietzia, Tsukamurella* and *Nocardioides*, in the manufacture of a medicament to for the treatment or prevention of melanoma and/or adenocarcinoma.

30

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera

Rhodococcus, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of an immune system imbalance in the elderly. Typically, an immune modulator composition or a pharmaceutical composition according to this aspect of the present invention may be an immune enhancer.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for enhancing the immune system which may result in for example, enhancement of growth or an increase in the efficiency of feed utilisation. Typically, the immune modulator composition or pharmaceutical composition according to this aspect of the present invention may be an immune enhancer. Advantageously, the immune modulator composition or pharmaceutical composition of the present invention may be used to replace antibiotics that are currently used to promote the growth of livestock. Suitably, the immune modulator composition of the present invention may be used either alone or in combination with other treatments. The term "livestock", as used herein refers to any farmed animal. Preferably, livestock is one or more of poultry, pigs (including piglets), sheep (including lambs), cows or bulls (including calves). More preferably, livestock means pigs – including piglets.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of an immune system imbalance in children.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*, in the manufacture of a medicament for the treatment or prevention of an adverse reaction to childhood vaccines – such as whooping cough vaccinations and the current MMR

vaccinations - and/or consequences thereof. The term "adverse reaction", as used herein, means a local or generalised disadvantageous response caused by or primed by the vaccine or the administration thereof, which typically occurs within a short time-frame but which can be delayed (for example by 6-months). An "adverse reaction" may include death of the child. The adverse reaction may be caused as a consequence of a separate event, the response to which has been negatively primed by the vaccine or the administration thereof.

The present invention further provides the use of an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* in the manufacture of a medicament for modifying a cellular immune response.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use as a medicament.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in or as a vaccine.

Suitably, the vaccine may be a prophylactic vaccine or a therapeutic vaccine.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use as an immune enhancer.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the

genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of one or more of an infection (e.g. a bacterial, viral, or parasitic infection, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, for example, major trauma stress, psychosocial stress and chronic stress, an allergy (e.g. asthma including allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses)), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (e.g. an immune system imbalance in children and the elderly).

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of a parasitic infection, such as, for example, one or more of malaria, trypanosomiasis, leishmaniasis and toxoplasmosis.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of stress, such as, for example, one or more of major trauma stress, psychosocial stress and chronic stress.

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of one or more of asthma (including allergic asthma), hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges for instance *Culicoides* (which causes Sweet Itch in horses). Preferably, the immune modulator composition or a pharmaceutical composition

comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* is for use in the treatment or prevention of asthma, including for example allergic asthma, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses).

5

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of one or more of heaves and/or COPD, particularly in horses.

10

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of cancer (for example melanoma and/or adenocarcinoma).

15

In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in the treatment or prevention of an immune system imbalance, particularly immunosenescence, in the elderly.

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In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* for use in enhancing the immune system which may result in for example, enhancement of growth or an increase in the efficiency of feed utilisation in, for example, livestock.

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In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the

genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* for use in the treatment or prevention of an immune system imbalance in children.

5 In a further aspect, the present invention provides an immune modulator composition or a pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* for use in the treatment or prevention of an adverse reaction to childhood vaccines – such as whooping cough vaccinations and the current MMR vaccinations – and/or consequences thereof.

10

In another aspect, the present invention provides the use of a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* in a vaccine or a medicament, wherein said whole cell of said bacterium modifies a cellular immune response.

15

In one aspect, the whole cell of the bacterium according to the present invention may downregulate a Th2 response.

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In another aspect, the whole cell of the bacterium according to the present invention may upregulate a Th1 response.

Suitably, the whole cell of the bacterium according to the present invention may downregulate a Th2 response and upregulate a Th1 response.

25 Alternatively, the whole cell of the bacterium according to the present invention may upregulate a Th1 response whilst not affecting a Th2 response.

Alternatively, the whole cell of the bacterium according to the present invention may downregulate a Th2 response, whilst also downregulating a Th1 response.

30

In another aspect, the present invention provides a method for treating or preventing a condition in a subject comprising administering an effective amount of a

pharmaceutical composition and/or immune modulator composition to a subject wherein the said composition modulates a cellular immune response.

Suitably the effective amount of the pharmaceutical composition and/or immune
5 modulator composition may be administered as a single dose. Alternatively, the effective amount of the pharmaceutical composition and/or immune modulator composition may be administered in multiple (repeat) doses, for example two or more, three or more, four or more, five or more, ten or more, or twenty or more repeat doses.

10 In a further aspect of the present invention, there is provided a method for protecting, including immunising, a subject comprising administering a pharmaceutical composition and/or immune modulator composition according to the present invention.

15 Preferably, a subject is protected, for example is immunised, against one or more of an infection (e.g. a bacterial, viral, or parasitic infection, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, for example, major trauma stress, psychosocial stress and chronic stress, an allergy (e.g. asthma including allergic
20 asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses)), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (e.g. an immune system imbalance in children and the elderly).

25

Preferably, a subject is immunised against one or more of malaria, trypanosomiasis, leishmaniasis and toxoplasmosis.

Preferably, a subject is protected against stress, such as, for example, one or more of major trauma stress, psychosocial stress and chronic stress.

30

Preferably, a subject is protected (including immunised) against one or more of asthma (including allergic asthma), hayfever, allergic dermatitis (eczema), anaphylactic shock,

allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges for instance *Culicoides* (which causes Sweet Itch in horses). More preferably, a subject is immunised against asthma including for example allergic asthma, and allergies to insect bites – such as midges, for instance

5 *Culicoides* (which causes Sweet Itch in horses).

Preferably, a subject is protected (including immunised) against one or more of COPD and/or heaves, particularly wherein the subject in a horse.

10 Preferably, a subject is protected against the development and/or the progression of a cancer, for example melanoma and/or adenocarcinoma.

Preferably, a subject is protected (including immunised) against an immune system imbalance in the elderly. In particular, the composition according to the present

15 invention may be used to regulate the subject's immune system.

Preferably, a subject is protected (including immunised) against an immune system imbalance in children. In particular, the composition according to the present invention may be used to regulate the subject's immune system, particularly the child's immune

20 system.

Preferably, a subject is protected (including immunised) against an adverse reaction to childhood vaccines and/or consequences thereof. In particular, the immune system of the subject is regulated, particularly a child's immune system, before and/or during

25 and/or after administration of the childhood vaccine.

The term "protected" as used herein means that the subject is less susceptible to the disease/disorder as compared with a subject not treated or administered with the compositions according to the present invention and/or that the subject is more able to

30 counter or overcome the disease/disorder as compared with a subject not treated or administered with the compositions according to the present invention.

In another aspect, the present invention provides administering an effective amount of a pharmaceutical composition and/or an immune modulator composition according to the present invention to a subject, wherein said composition is co-administered with an antigen or antigenic determinant.

5

When the composition is co-administered with an antigen or antigenic determinant in accordance with the present invention the antigen or antigenic determinant may suitably be an antigen or antigenic determinant from one or more of the following: BCG (bacillus of Calmette and Guerin) vaccine, diphtheria toxoid vaccine, diphtheria/tetanus/pertussis (DTP or Triple) vaccine, pertussis vaccine, tetanus toxoid vaccine, measles vaccine, mumps vaccine, rubella vaccine, OPV (oral poliomyelitis vaccine) and *Mycobacterium vaccae*, or part thereof (as taught in GB0025694.1). Suitably two or more, or three or more, of such antigens or antigenic determinants may be co-administered with a pharmaceutical composition or an immune modulator composition according to the present invention.

15

Preferably, a medicament according to the present invention is used for the treatment or prevention of one or more of an infection (e.g. a bacterial, viral, or parasitic infection, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, for example, major trauma stress, psychosocial stress and chronic stress, an allergy (e.g. asthma including allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses)), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (e.g. an immune system imbalance in children and the elderly).

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Preferably, a medicament according to the present invention is used for the treatment or prevention of one or more of malaria, trypanosomiasis, leishmaniasis and toxoplasmosis.

30

Preferably, a medicament according to the present invention is used for the treatment or prevention of stress, such as, for example, major trauma stress, psychosocial stress and chronic stress.

- 5 Preferably, a medicament according to the present invention is used for the treatment or prevention of one or more of asthma (including allergic asthma), hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses). More preferably, a
10 medicament according to the present invention is used for the treatment or prevention of asthma, including allergic asthma, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses).

- Preferably, a medicament according to the present invention is used for the treatment
15 or prevention of one or more of COPD and heaves.

Preferably, a medicament according to the present invention is used for the treatment or prevention of cancers (for example melanoma and/or adenocarcinoma).

- 20 Preferably, a medicament according to the present invention is used for the treatment or prevention of an immune system imbalance in the elderly.

Preferably, a medicament according to the present invention is used for the treatment or prevention of an immune system imbalance in children.

25

Preferably, a medicament according to the present invention is used for the treatment or prevention of an adverse reaction to childhood vaccines and/or consequences thereof.

- 30 In a further aspect of the present invention, a pharmaceutical composition or an immune modulator composition according to the present invention may comprise bacteria from more than one of the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*,

Tsukamurella and *Nocardioides*. Suitably, the composition may comprise two or more, or three or more, bacteria from any one of the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*.

5 Preferably, the bacteria for use in accordance with the present invention are from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* and include *Gordonia bronchialis*, *Nocardia asteroides*, *Dietzia maris*, *Tsukamurella paurometabola*, *Rhodococcus ruber*, *Nocardioides albus* and *Tsukamurella inchonensis*. Suitably, the species used from each particular genus are ones which can
10 be grown on medium, which is a low, preferably non-, antigenic medium. By way of example only, a suitable non-antigenic medium is Sauton's medium.

More preferably, the bacteria for use in accordance with the present invention are from the genus *Rhodococcus*, including *Rhodococcus ruber* (previously known as *Nocardia rubra*),
15 *Rhodococcus rhodocrous*, *Rhodococcus rhodnii*, *Rhodococcus coprophilus*, *Rhodococcus opacus*, *Rhodococcus erythropolis*.

More preferably, the bacteria for use in accordance with the present invention is
20 *Rhodococcus ruber*.

Preferably, the bacterium according to the present invention is killed prior to use. Preferably, the bacterium according to the present invention is killed by heat-treatment thereof, for example, heat-treatment in an autoclave at 121°C for 15 minutes. Other
25 suitable treatments for killing the bacterium may include ultraviolet or ionising radiation or treatment with chemicals such as phenol, alcohol or formalin.

Preferably, the bacterium according to the present invention is purified and/or isolated.

Preferably, the bacterium according to the present invention is suspended in water or
30 buffered saline, suitably borate buffered at pH 8.

The term "subject", as used herein, means an animal. Preferably, the subject is a mammal, including for example livestock and humans. In some aspects of the present invention, the subject may suitably be a human.

- 5 The term "immune modulator" as used herein includes a vaccine.

THERAPEUTIC USES

- 10 The immune modulators of the present invention may be used in therapy. In particular such compounds may be used to modulate T lymphocyte responses *in vivo* and/or other cells involved in an immune response *in vivo*.

- 15 Immune modulator/pharmaceutical compositions capable of modulating, in particular blocking, T cell proliferation and/or differentiation and/or activity may be used against any disorder which is susceptible to prevention or treatment by the modulation of an adaptive immune response, i.e. a cellular immune response.

- 20 Suitably, the compositions according to the present invention are used to modulate a cellular immune response to treat or prevent one or more of an infectious disease (such as a bacterial infection eg. methycillin-resistant *Staphylococcus aureus*, tuberculosis, including multidrug resistant tuberculosis and leprosy; chronic viral infections, for example hepatitis and HIV; latent viral infections, such as shingles (*Herpes zoster*) or cold sores (*Herpes simplex*) for example; or parasitic infections, for example malaria, trypanosomiasis, leishmaniasis and toxoplasmosis), an allergy, such as allergic
25 dermatitis or allergic asthma, stress – such as major trauma stress, psychosocial stress and chronic stress, and cancer (for example, the composition may be administered regularly throughout adult life to counter the effects of tobacco).

- 30 A more extensive list of disorders is given in WO-A-98/09985. For ease of reference, part of that list is now provided: inflammation associated with hypersensitivity, allergic reactions, asthma, Raynaud's phenomenon, inflammation associated with aphthous ulceration, ulcerative colitis, hepatic fibrosis, liver cirrhosis or other hepatic

diseases, dermatitis, in particular atopic dermatitis e.g. eczema, periodontal diseases or other dental diseases, to treat or ameliorate monocyte or leukocyte proliferative diseases, e.g. leukaemia, other cancers.

5 Infectious diseases

Compositions capable of modulating, in particular stimulating (i.e. inducing or enhancing) T cell proliferation and/or differentiation or of preventing the induction of or reversing T cell anergy may be used generally to boost or induce T cell immune responses. Virtually all adaptive immune responses require the activation of T cells and their differentiation into cytokine-producing cells. Thus, these compositions may be used generally to prevent and/or treat infectious diseases - such as viral, bacterial. Suitably, these compositions may be used to prevent and treat parasitic infections, e.g. malaria, leishmaniasis, toxoplasmosis and trypanosomiasis).

15

In one aspect of the present invention the infection is preferably trypanosomiasis.

Allergy

20 The compositions of the present invention may also be used to prevent and treat allergies (eg. asthma including allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings - such as nettle and insect stings, and allergies to insect bites - such as midges for instance *Culicoides* (which causes Sweet Itch in horses).

25

Sweet Itch is one of the commonest skin diseases seen in horses, particularly in wild horses and/or ponies. About 3% of horses in the U.K. are affected to some degree. Most horses show signs between 1 and 4 years of age and the condition generally worsens during summer. Certain breeds are particularly prone to the disease. Shires, Hackneys and Welsh ponies have all been suggested as susceptible breeds. Sweet Itch is caused by hypersensitivity to the bites of the tiny fly *Culicoides*. In the UK, the fly is present from April to October but peaks in numbers in May to September. The flies

30

feed on the horse at specific sites usually around the tail head and under the mane. There are 20 species of *Culicoides* present in the UK and some feed underneath the horses' abdomen.

- 5 By way of example only, the compositions of the present invention may also be used to prevent and/or treat anaphylactic shock by administering a subject, which subject has a predisposition to suffer from anaphylactic shock (for instance a subject who is known to have an allergy, such as an allergy to peanuts for instance) with a composition according to the present invention, thus to reduce the subject's
10 predisposition to anaphylactic shock should they (accidentally) come into contact with the antigen, e.g. peanuts for instance, towards which they may react adversely.

Heaves/COPD

- 15 The compositions of the present invention may also be used to prevent and treat

Heaves is an equine lung disease with similarities to human asthma and COPD (Chronic Obstructive Pulmonary Disease). The clinical signs in the horse are initiated by an allergic response to the particles in hay dust in lungs already damaged with a
20 degree of fibrosis. It is most often seen in older horses (greater than six years old) that are stabled during the winter months. Hay contains microorganisms - such as bacteria and fungi as well as tiny particles of feed grains, plants, faeces, dander, and pollen. These tiny particles become aerosolised in hay dust and elicit an allergic response and fibrosis when they are inhaled by horses with heaves. The primary microorganisms
25 believed to be involved in the etiology of heaves are *Aspergillus fumigatus*, *Thermoactinomyces vulgaris* and *Faenia rectivirgula*. Both reduction of the bronchospasm of asthma and the fibrosis of COPD are within the scope of the patent.

Stress

30

Stress is often presented as a symptom of modern living, the high pressure executive lifestyle, the consequences of which are widely perceived as leading to major

pathological conditions such as gastric ulcers, hypertension, heart disease and strokes. Other major stressful events in life such as divorce, bereavement and moving house are seen as high risk factors for heart disease. These are not misconceptions, the farming industry is well aware of the economic losses resulting from subjecting

5 livestock to major stresses such as overcrowding, confinement and transportation leading to an increased susceptibility to infection and the precipitation of underlying pathology. Research by doctors and scientists is producing an increasing volume of published work showing definable stresses such as confinement can result in significant changes in endocrine (hormone) activity which subsequently can affect the

10 body's immune functions. This can be noticeably demonstrated in major trauma stress (including surgical stress) in which the cell mediated immune response is dramatically paralysed Faist (1996).

Elenkov IJ (1999) report recent evidence indicating that glucocorticoids and

15 catecholamines, the end products of the stress system, and histamine, a product of activated mast cells, might selectively suppress cellular immunity, and favour humoral immune responses. This is mediated by a differential effect of stress hormones and histamine, on Th1/Th2 patterns and type 1/type 2-cytokine production. Thus, systemically, stress might induce a Th2 shift, while, locally, under certain conditions,

20 it might induce pro-inflammatory activities through neural activation of the peripheral corticotrophin-releasing factor-mast cell-histamine axis.

Paik (2000) and Kay (2001) in independent studies of academic stress, examined the immunological profiles of students during non exam and exam periods. They report a

25 significant reduction in IL-2 and interferon gamma production and an increase in IL-6. This indicates that the body's immune system responds to stressful episodes by a down regulation of Th-1 cytokines and a selective up-regulation of the Th2 cytokines.

Iwakabe (1998), using a mouse model of restraint stress, reports the skewing of the

30 immune response towards Th2 dominant immunity.

This stress hormone induced switch towards Th2 immune imbalance is also reported in non major, chronic stress situations such as psychosocial stress amongst workers overwintering at the Australian National Antarctic Research Expedition stations, (Mehta (2000)). They also report an associated increase in latent virus reactivations.

5

Similar stress hormone and immunological changes are reported from chronic stress in care givers of dementia patients (Bauer (2000)) and in astronauts during the Euromir 95 mission (Norbiato (1998)). Of particular concern was the astronauts increased susceptibility to infection.

10

The body is designed to recover from stress and in acute stress clearly does as the risk of infection recedes with the patient's recovery from the major trauma.

Chronic stress however appears to maintain the Th2 dominated immune imbalance.

15

This is a very serious consequence as all of the quoted authors allude to stress through the above mechanisms, possibly influencing the onset and/or course of infectious, inflammatory, allergic and neoplastic diseases.

This consequence is further supported by Lawrence (2000).

20

An immunoregulator, preferably, an orally administered immunoregulator, according to the present invention, which stimulates the Th1 response and down regulates Th2 may restore the healthy balance of the immune system and thus reduce the increased risk of serious illness associated with chronic stress.

25

Preferably, the composition according to the present invention is used to treat and/or prevent stress, in particular major trauma stress, psychosocial stress and chronic stress.

Immune system imbalance

30

An immune system imbalance - such as an upregulation, downregulation or inappropriately regulated cellular immune response - may occur at any time in the life

of a subject. Suitably, the compositions may be used to modulate an immune system imbalance. That is to say, the compositions according to the present invention may be used to treat and/or prevent an immune system imbalance.

5 (a) In children

Suitably, the immune modulator composition or a pharmaceutical composition may be used to modulate an immune system imbalance, in children, including babies, infants and juveniles. An immune system imbalance - such as an upregulation,
10 downregulation or inappropriately regulated cellular immune response - may occur in children following vaccination, for example following childhood vaccinations. Such an immune system imbalance may result in conditions such as the onset of allergies, i.e. allergic dermatitis and allergic asthma.

15 With the aim of protecting children from infections, repeated injections against Diptheria, Tetanus, Pertussis, Polio, Measles, Mumps and the Rubella are given. All of these are judged necessary and pressure is exerted by Health authorities to ensure that children are presented for vaccination at the appropriate time. However, most vaccinations given in early life contain an alum adjuvant, which has important
20 immunological consequences. Alum is a potent stimulus to the Th2 pattern of response and the consequential immune dysregulation causes the child to become vulnerable to the development of allergies and possibly cancer for example.

It is possible to re-educate the immune system to a proper recognition, regulation and
25 response both the self and to the rest of the world.

Suitably, the immune modulator composition or a pharmaceutical composition may also be used for the treatment or prevention of an adverse reaction to childhood vaccines – such as whooping cough vaccinations and the current MMR vaccinations - and/or consequences thereof.

(b) Immune system imbalance in the elderly

An immune system imbalance - such as an upregulation, downregulation or inappropriately regulated cellular immune response, in particular downregulation, for example a deterioration of immune function – may occur in older people, generally in excess of 60 years. In elderly people, a downregulation in the cellular immune response is generally referred to as immunosenescence. Typically, the deterioration of immune function may lead to increased susceptibility to infectious diseases and neoplasia for example. The number of old people as a proportion of the population is dramatically increasing and geriatric medicine is becoming an important aspect of clinical practice. It is not surprising therefore that research has focused on the mechanisms of immunosenescence and the links between the health of the immune system and longevity. Goronzy (2001), examined the varying efficacy of influenza vaccination in the elderly. In this study, only 17% of subjects showed a rise in titre to all 3 haemagglutinins (successful vaccination) 1 month following vaccination and that 46% showed no demonstrable response at all. It was proposed that responsiveness to influenza vaccination is a useful biological marker of immunosenescence. A number of researchers have studied various aspects of the immune function in the elderly. For example, Lio (2000) studied cytokine responses, Solana (2000) studied NK and NKt cells, and Ginaldi (1999) suggested that a Th1 to Th2 cytokine production shift and an increased production of proinflammatory cytokines could explain many aspects of age-associated pathological events, such as atherosclerosis and osteoporosis. Accordingly, a non-pathological stimulation of the immune system which drives the cytokine response away from the proinflammatory Th2 towards Th1 is required. Preferably, such an immune modulator reduces the mortality from acute infection, counters the onset and reduces the morbidity of age related autoimmune

disease and possibly reduces the rate of neoplastic disease, all of which are associated with immunosenescence.

Typically, an immune modulator composition or a pharmaceutical composition
5 according to this aspect of the present invention may be a immune enhancer.

The potential role of probiotic commensal gut or dairy bacteria have been investigated in this area. For example, milk products supplemented with 5×10^9 or 5×10^{10} *Bifidobacterium lactis* or *Lactobacillus rhamnosus* per dose taken daily for 3 weeks
10 has been reported to increase the numbers of peripheral blood NK, CD4 and CD25 cells and generally boost systemic cellular immunity in the elderly (Gill, (2001)).

Currently a number of products containing high numbers of Lactobacilli and other intestinal commensal flora are being actively promoted as "lifestyle enhancers". A
15 review by Sanders (2001), on the claimed probiotic effects of *Lactobacillus acidophilus*, available as a drug since 1950, suggests its effects require further validation and clarification of the mechanism of action. Whilst there is a beneficial effect from replacing the bowel flora after diarrhoeal disease and combating candidiasis following antibiotic therapy, immune stimulation appears unreliable and
20 short lived. However, this work clearly identifies a role for a potent orally administered immune modulator, preferably killed so as to avoid the difficulties of maintaining live products.

Oral vaccination is a long established successful mechanism for inducing local
25 protective immunity against oral/faecal pathogens - such as polio. However, orally administered vaccines have also been shown to evoke systemic protective immune responses both cell mediated and humoral. Sharpe (2002), used an orally administered adenovirus construct containing measles virus antigen to induce systemic antibody and splenic lymphocyte responses to the measles antigen. Manube (2002) has developed a
30 model to show that orally administered attenuated *Mycobacterium microti* provides a higher level of protection to an aerosol challenge with tuberculosis than traditional subcutaneously administered BCG.

Kim (2001) showed that feeding with Japanese cedar pollen produced oral tolerance to specific allergy induced by subsequent injection of pollen in oil. This was associated with decreased specific immunoglobulin levels and a significant reduction in interleukin-4 production *ie* the TH2 response.

Therefore, a systemic immune response may be both stimulated and modulated by administration – such as oral administration - of a suitable immune modulator.

Suitably, in one aspect of the present invention it is envisaged that a whole cell of the bacterium according to the present invention, may be included in food preparations and/or may be supplied as a type of “remedy”, preferably orally.

Enhancing the immune system

The compositions of the present invention may be used in the manufacture of a medicament for enhancing the immune system in an animal , preferably a mammal, more preferably livestock, which may result in for example, enhancement (e.g. promotion) of growth and/or an increase in the efficiency of feed utilisation and/or generally increased well-being (i.e. the overall health of the subject is improved) in the subject. Typically, the immune modulator composition or pharmaceutical composition according to this aspect of the present invention may be an immune enhancer.

Advantageously, the immune modulator composition or pharmaceutical composition of the present invention may be used to replace antibiotics that are currently used to promote the growth of livestock.

The present invention also contemplates the genera of the present invention being administered in combination with known probiotic bacteria, for modification of the cellular immune response.

Commercially at present antibiotics are commonly used as dietary enhancing feed additives (or growth promoters) and are incorporated into animal feed. However, the EU, is expected to introduce a complete ban on the non-clinical use of antibiotics in animal husbandry. Therefore, the market requires effective alternatives.

5

The immune modulator compositions or pharmaceutical compositions of the present invention may be administered as a food additive when used to enhance the immune system.

10 Cancer

Suitably, the compositions according to the present invention are used to modulate a cellular immune response to treat and/or prevent cancer. In particular it is envisaged that the compositions according to the present invention may be used to protect a subject against developing and/or the progression of a cancer. In particular, the subject with a modulated cellular immune response may be less susceptible to the development of cancer.

15

In particular, during cancer growth an unregulated increase in Th2 is observed.

20

Cancer is a disease that affects many people, with 65 per cent of cases occurring in those over 65. As the average life expectancy in the UK has almost doubled since the mid-nineteenth century, the population at risk of cancer has grown. Death rates from other causes of death, such as heart disease, have fallen in recent years while deaths from cancer have remained relatively stable. The result is that 1 in 3 people will be diagnosed with cancer during their lifetime and 1 in 4 people will die from cancer.

25

Examples of cancer include bladder, brain tumor, breast cancer, cervical cancer, colon and rectal cancer, adenocarcinoma, endometrial cancer, esophageal cancer, kidney cancer, leukaemia, liver cancer, lung cancer, melanoma, myeloma, non-hodgkin's lymphoma, ovarian cancer, pancreatic cancer, prostate cancer, sarcoma, soft tissue and stomach cancer.

30

In addition, persistent smoking of tobacco, and to a lesser, extent passive smoking, has been associated with carcinomas of the parts directly in contact with smoke, oropharynx, trachea, lungs, oesophagus and stomach. As well as these, distant
5 tumours such as those of the kidney, bladder, pancreas, liver and myeloid leukaemia may be increase by smoking of tobacco. In the present invention, it is envisaged that compositions according to the present invention could be administered to smokers of tobacco in an attempt to reduce the smokers' risk of developing carcinomas associated with tobacco smoking.

10

Suitably, the cancer may be an adenocarcinoma or a melanoma.

T HELPER CELLS

15 The term 'Th1' as used herein refers to a type 1 T-helper cell (Th1). The term may also be used herein to refer to the response mediated by or through such a cell type. Such a response may include one or more of the secretion of Interleukin-2 (IL-2), the secretion of Interferon-gamma (IFN- γ), activation of macrophages, activation of cytotoxic T-cells, or any other Th1-associated event. Thus, the term 'Th1' may
20 include Th1 cell(s) as well as the immune response(s) which such cell(s) produce.

The term 'Th2' as used herein refers to a type 2 T-helper cell (Th2). The term may also be used herein to refer to the response mediated by or through such a cell type. Such a response may include one or more of the secretion of Interleukin-4 (IL-4), the
25 secretion of the splice variant interleukin IL-482, the secretion of Interleukin-5 (IL-5), increase in levels of cell determinant 30 (CD30) on lymphocytes, increase in levels of Immunoglobulin-E (IgE) in the blood or eosinophils in the blood, or any other Th2-associated event. Thus, the term 'Th2' may include Th2 cell(s) as well as the immune
30 response(s) which such cell(s) produce.

30

It is known that various conditions may result in or from an unregulated or inappropriately regulated cellular immune response, in particular in the activation

and/or proliferation of Th1 and/or Th2, which if left unregulated or inappropriately regulated has been found to result in one or more detrimental effects on the subject.

In particular, such an unregulated or inappropriately regulated cellular immune response has been found to occur following vaccination, eg. following childhood vaccinations, and is thought to result in conditions such as the onset of allergies, i.e. allergic dermatitis and allergic asthma. By way of example, Lewis D *Curr Opin Immunol* 2002; 14: 644 report that Th2 immune responses mediated by the secretion of IL-4, IL-5 and IL-13 are key in the pathogenesis of atopic disorders, including allergen-induced asthma, rhinoconjunctivitis and anaphylaxis. Although such responses are downregulated to some degree by conventional specific immunotherapy, this approach is only partially effective and has a substantial risk of adverse effects. Many strategies for immunotherapeutic prophylaxis and for treatment of atopic diseases have been devised on the basis of mouse allergy models, including the downregulation of Th2 responses by the induction of regulatory T cell activity, Th2 to Th1 immune deviation, Th1 crossregulation of Th2 immune responses, anergy and immunosuppressive cytokines. Choi & Koh *Ann Allergy Asthma Immunol* 2002; 88: 584-91 examined whether BCG vaccination of adult patients with asthma, a Th2-associated allergic disease, is clinically effective. It was shown that BCG vaccination improved lung function and reduced medication use in adults with moderate-to-severe asthma. This amelioration was accompanied by a suppressed Th2-type immune response, suggesting that BCG vaccination might be an effective therapeutic modality against asthma. von Hertzen *J Allergy Clin Immunol* 2002; 109: 923-8 outlined the possibility that prolonged maternal stress associated with sustained excessive cortisol secretion could affect the developing immune system - especially Th1/Th2 cell differentiation which may further increase the susceptibility to asthma and atopy in genetically predisposed individuals.

In addition, an unregulated or inappropriately regulated cellular immune response has been observed during disease progression. In particular during cancer growth an unregulated increase in Th2 is observed. By way of example, Maraveyas *et al. Ann Oncol* 1999; 10: 817-24 have studied the efficacy of SRL 172 vaccine in patients with

cancer ie. advanced stage IV (AJCC) malignant melanoma. Induction of intracellular cytokines (IL-2 and INF-gamma) in peripheral blood lymphocytes (PBLs) from these patients was assayed and correlated to clinical outcome. It was demonstrated that SRL 172 was effective in inducing intracellular IL-2 responses in a significant number of patients with stage IV (AJCC) melanoma. Stanford *et al. International Journal of Pharmaceutical Medicine* 1999; 13: 191-195 report that there is increasing evidence that effective anti-tumour immune responses are likely to be mediated by type 1 cytokines. Recent investigations indicate that heat-killed *Mycobacterium vaccae*, is a reliable Th1 adjuvant and preliminary clinical trials indicate beneficial effects in melanoma, and cancer of the prostate and lung. More extensive controlled studies are currently being conducted to confirm these findings.

An unregulated or inappropriately regulated cellular immune response has also been observed during infection and particularly chronic infection, for example during progressive tuberculosis, lepromatous leprosy, visceral leishmaniasis and HIV infection, during allergies and during Raynaud's disease. By way of example, Clerici & Shearer *GM Immunol Today* 1993; 14: 107-11 propose that a Th1 to Th2 switch is a critical step in the etiology of HIV infection. Clerici & Shearer *Immunol Lett* 1996; 51: 69-73 show that HIV-specific cell mediated immunity may be the main correlator of protection against HIV infection and against the progression of HIV infection to AIDS. Abbot NC *et al. European Journal of Vascular and Endovascular Surgery* 2002 (in press) evaluated immunotherapy as a means of improving peripheral blood flow in chronic leprosy patients by administration of heat-killed *Mycobacterium vaccae*. It was shown that immunotherapy, given 18 months earlier, significantly improved blood flow and temperature sensation, in fully-treated, chronic, leprosy patients.

Accordingly, an aim of the present invention is to promote and establish the regulation of a cellular immune response, including the regulation or modulation of Th1 and/or Th2, in such a way so as to overcome the negative effects of the unregulated or inappropriately regulated cellular immune response.

Suitably, the use of an immune modulator composition and/or pharmaceutical composition according to the present invention modulates the Th1 or Th2 response, i.e. a Th1 or Th2 response that results in, for example, tissue damage.

- 5 Suitably, the use of an immune modulator composition and/or pharmaceutical composition according to the present invention may decrease the Th1 response and decrease the Th2 response. By way of example, such an immune modulator and/or pharmaceutical composition may be useful in the treatment of diabetes for example.
- 10 Suitably, the use of an immune modulator composition and/or pharmaceutical composition according to the present invention may increase the Th1 response without affecting the Th2 response. By way of example, such an immune modulator and/or pharmaceutical composition may be useful as an immune enhancer.
- 15 Suitably, the use of an immune modulator composition and/or pharmaceutical composition according to the present invention may increase the Th1 response and decrease the Th2 response. By way of example, such an immune modulator and/or pharmaceutical composition may be useful in the treatment of asthma.
- 20 Suitably, a skilled person can test a specific species of each genus according to the present invention to determine its specific Th1/Th2 response.

An unregulated or inappropriately regulated immune response may play a role in the establishment of disease due to the fact that some diseases cause shifted Th1 and/or
25 Th2 responses. Accompanying these atypical Th1 and Th2 reactions are a series of abnormal inflammatory responses, which may take part in the mechanisms underlying tissue pathology.

By way of example only, the immune modulator composition and/or pharmaceutical
30 composition according to the present invention may counteract the disadvantages of reduced contact with environmental influences (for example, antigens) commensurate with modern life, may counteract the influence of treatment of an infection (eg. a

parasitic infection, such as, for example, malaria, trypanosomiasis, leishmaniasis, and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, such as, for example, major trauma stress, psychosocial stress and chronic stress, an allergy (eg. asthma including asthma, allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and insect stings, and allergies to insect bites – such as midges for instance *Culicoides* (which causes Sweet Itch in horses), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (eg. an immune system imbalance in children eg. the undesirable effect of childhood vaccines and the elderly).

VACCINES

The preparation of vaccines which contain one or more substances as an active ingredient(s), is known to one skilled in the art. Typically, such vaccines are prepared as injectables, either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid prior to injection may also be prepared. The preparation may also be emulsified, or the active ingredient(s) encapsulated in liposomes. The active ingredients are often mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredient. Suitable excipients are, for example, water, saline, dextrose, glycerol, ethanol, or the like and combinations thereof. Alternatively, the vaccine may be prepared, for example, to be orally ingested and/or capable of inhalation.

In addition, if desired, the vaccine may contain minor amounts of auxiliary substances such as wetting or emulsifying agents and pH buffering agents.

ADMINISTRATION

Typically, a physician will determine the actual dosage of a vaccine, immune modulator composition and pharmaceutical composition which will be most suitable for an individual subject and it will vary with the age, weight and response of the

particular patient. The dosages below are exemplary of the average case. There can, of course, be individual instances where higher or lower dosage ranges are merited.

Preferably, the actual dosage that is used results in minimal toxicity to the subject.

5

The compositions of the present invention may be administered by direct injection. The composition may be formulated for parenteral, mucosal, intramuscular, intravenous, subcutaneous, intraocular, intradermal or transdermal administration. Suitably, the composition according to the present invention may be administered at a
10 dose of $10^6 - 10^9$ organisms. Typically, the composition according to the present invention may be administered at a dose of $10^8 - 10^9$ bacteria for human and animal use.

If the compositions of the present invention are to be administered as immune enhancers, then $10^6 - 10^9$ organisms per dose, preferably, $10^8 - 10^9$ bacteria per dose for
15 human and animal use may be administered at regular intervals.

The term "administered" includes delivery by delivery mechanisms including injection, lipid mediated transfection, liposomes, immunoliposomes, lipofectin, cationic facial amphiphiles (CFAs) and combinations thereof, or even viral delivery. The routes for
20 such delivery mechanisms include but are not limited to mucosal, nasal, oral, parenteral, gastrointestinal, topical, or sublingual routes.

The term "administered" includes but is not limited to delivery by a mucosal route, for example, as a nasal spray or aerosol for inhalation or as an ingestible solution; a
25 parenteral route where delivery is by an injectable form, such as, for example, an intravenous, intramuscular, intradermal or subcutaneous route.

The term "co-administered" means that the site and time of administration of each of the adjuvants(s), antigen(s) and/or antigenic determinant(s) of the present invention are
30 such that the necessary modulation of the immune system is achieved. Thus, whilst the antigen(s) and adjuvant(s) may be administered at the same moment in time and at the same site, there may be advantages in administering the antigen(s) and/or antigenic

determinant(s) at a different time and to a different site from the adjuvant(s). The antigen(s) and/or antigenic determinant(s) and adjuvant(s) may even be delivered in the same delivery vehicle - and the antigen(s) and/or antigenic determinant(s) and adjuvant(s) may be coupled and/or uncoupled and/or genetically coupled and/or uncoupled. By way of example only, the immune modulator composition according to the present invention may be administered before, at the same time or post administration of one or more antigens or further antigens.

The antigen, antigenic determinant, peptide or homologue or mimetic thereof may be administered separately or co-administered to the host subject as a single dose or in multiple doses.

The immune modulator composition and/or pharmaceutical composition of the invention may be administered by a number of different routes such as injection (which includes parenteral, subcutaneous, intradermal and intramuscular injection) intranasal, mucosal, oral, intra-vaginal, urethral or ocular administration.

Preferably, in the present invention, administration is by injection. More preferably the injection is intradermal.

Preferably, in the present invention, administration is by an orally acceptable composition.

For vaccination the composition can be provided in 0.1 to 0.2 ml of aqueous solution, preferably buffered physiological saline, and administered parenterally, for example by intradermal inoculation. The vaccine according to the invention is preferably injected intradermally. Slight swelling and redness, sometimes also itching may be found at the injection site. The mode of administration, the dose and the number of administrations can be optimised by those skilled in the art in a known manner.

ANTIGENS

As used herein, an "antigen" means an entity which, when introduced into an immunocompetent host, modifies the production of a specific antibody or antibodies that can combine with the entity, and/or modifies the relevant Th response, such as Th2 and/or Th1. The antigen may be a pure substance, a mixture of substances or soluble or particulate material (including cells or cell fragments or cell sonicate). In this sense, the term includes any suitable antigenic determinant, cross reacting antigen, alloantigen, xenoantigen, tolerogen, allergen, hapten, and immunogen, or parts thereof, as well as any combination thereof, and these terms are used interchangeably throughout the text.

The term "antigenic determinant or epitope" as used herein refers to a site on an antigen which is recognised by an antibody or T-cell receptor, or is responsible for evoking the T-helper cell response. Preferably it is a short peptide derived from or as part of a protein antigen. However the term is also intended to include glycopeptides and carbohydrate epitopes. The term also includes modified sequences of amino acids or carbohydrates which stimulate responses which recognise the whole organism.

It is advantageous if the antigenic determinant is an antigenic determinant of the infectious agent which causes the infectious disease.

A "preventative" or "prophylactic" vaccine is a vaccine which is administered to naive individuals to prevent development of a condition, such as by stimulating protective immunity.

A "therapeutic" vaccine is a vaccine which is administered to individuals with an existing condition to reduce or minimise the condition or to abrogate the immunopathological consequences of the condition.

ADJUVANTS

The term 'adjuvant' as used herein means an entity capable of augmenting or participating in the influencing of an immune response. An adjuvant is any substance
5 or mixture of substances that assists, increases, downregulates, modifies or diversifies the immune response to an antigen..

The immune modulator composition and/or pharmaceutical composition according to the present invention may comprise one or more adjuvants which enhance the
10 effectiveness of the immune modulator composition and/or pharmaceutical compositions. Examples of additional adjuvants which, may be effective include but are not limited to: aluminium hydroxide, aluminium phosphate, aluminium potassium sulphate (alum), beryllium sulphate, silica, kaolin, carbon, water-in-oil emulsions, oil-in-water emulsions, muramyl dipeptide, bacterial endotoxin, lipid X, *Corynebacterium*
15 *parvum* (*Propionobacterium acnes*), *Bordetella pertussis*, *Mycobacterium vaccae*, polyribonucleotides, sodium alginate, lanolin, lysolecithin, vitamin A, interleukins such as interleukin 2 and interleukin-12, saponin, liposomes, levamisole, DEAE-dextran, blocked copolymers or other synthetic adjuvants. Such adjuvants are available commercially from various sources, for example, Merck Adjuvant 65 (Merck and
20 Company, Inc., Rahway, N.J.) or Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, Michigan). Only aluminium hydroxide is approved for human use. Some of the other adjuvants, such as *M. vaccae* for example, have been approved for clinical trials.

25 Suitably, the adjuvant may be a whole cell of a bacterium from any one of the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides*.

In the art, it is known that DNA vaccines, which are essentially DNA sequences attached to gold particles and which are fired into the skin by a helium gun, are
30 efficient vaccine delivery systems. Unlike conventional vaccines, these DNA vaccines do not require a traditional adjuvant component. In accordance with a further aspect of the present invention, the immune modulator composition as defined herein may

suitably be used in conjunction with such DNA vaccines to augment or participate in the influencing of the immune response.

PHARMACEUTICAL COMPOSITIONS

5

The present invention also provides a pharmaceutical composition comprising a therapeutically effective amount of a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* and optionally a pharmaceutically acceptable carrier, diluent or excipients (including combinations thereof).

10

The pharmaceutical composition may comprise two components – a first component comprising an antigen and a second component comprising an adjuvant thereof. The first and second component may be delivered sequentially, simultaneously or together, and even by different administration routes.

15

The pharmaceutical compositions may be for human or animal usage in human and veterinary medicine and will typically comprise any one or more of a pharmaceutically acceptable diluent, carrier, or excipient. Acceptable carriers or diluents for therapeutic use are well known in the pharmaceutical art, and are described, for example, in Remington's Pharmaceutical Sciences, Mack Publishing Co. (A. R. Gennaro edit. 1985). The choice of pharmaceutical carrier, excipient or diluent can be selected with regard to the intended route of administration and standard pharmaceutical practice. The pharmaceutical compositions may comprise as - or in addition to - the carrier, excipient or diluent any suitable binder(s), lubricant(s), suspending agent(s), coating agent(s), solubilising agent(s).

20

25

Preservatives, stabilisers, dyes and even flavouring agents may be provided in the pharmaceutical composition. Examples of preservatives include sodium benzoate, sorbic acid and esters of p-hydroxybenzoic acid. Antioxidants and suspending agents may be also used.

30

There may be different composition/formulation requirements dependent on the different delivery systems. By way of example, the pharmaceutical composition of the present invention may be formulated to be delivered using a mini-pump or by a mucosal route, for example, as a nasal spray or aerosol for inhalation or ingestable solution, or parenterally in which the composition is formulated by an injectable form, for delivery, by, for example, an intravenous, intramuscular, intradermal or subcutaneous route. Alternatively, the formulation may be designed to be delivered by both routes.

10 Preferably in the present invention the formulation is of injectable form. More preferably the formulation is intradermally injected.

Preferably in the present invention the formulation is an orally acceptable composition.

15 Where the agent is to be delivered mucosally through the gastrointestinal mucosa, it should be able to remain stable during transit through the gastrointestinal tract; for example, it should be resistant to proteolytic degradation, stable at acid pH and resistant to the detergent effects of bile.

20 Where appropriate, the pharmaceutical compositions can be administered by inhalation, in the form of a suppository or pessary, topically in the form of a lotion, solution, cream, ointment or dusting powder, by use of a skin patch, orally in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs, solutions or
25 suspensions containing flavouring or colouring agents, or they can be injected parenterally, for example intravenously, intramuscularly, intradermally or subcutaneously. For parenteral administration, the compositions may be best used in the form of a sterile aqueous solution which may contain other substances, for example enough salts or monosaccharides to make the solution isotonic with blood. For buccal
30 or sublingual administration the compositions may be administered in the form of tablets or lozenges which can be formulated in a conventional manner.

PHARMACEUTICAL COMBINATIONS

- The agent of the present invention may be administered with one or more other pharmaceutically active substances. By way of example, the present invention covers
- 5 the simultaneous, or sequential treatments with an immune modulator composition and/or pharmaceutical composition according to the present invention, and one or more steroids, analgesics, antivirals, interleukins such as IL-2, or other pharmaceutically active substance(s).
- 10 It will be understood that these regimes include the administration of the substances sequentially, simultaneously or together.

IMMUNE ENHANCER

- 15 The term "immune enhancer" as used herein means one or more bacteria either isolated or in culture which when administered to a subject benefit the health of that subject. Preferably, this benefit is achieved by the modification of the cellular immune response of the subject.
- 20 In accordance with the present invention, immune enhancers may be used, for example, for the treatment or prevention of an immune system imbalance in a subject, preferably a child or an elderly subject, or for enhancing the immune system of a subject, for example of a mammal, particularly of livestock or of humans.
- 25 The immune enhancers may be administered by consumption in specially designed food or in animal feeds, for example pig animal feeds supplemented with the bacteria of the present invention.

- The immune enhancers may also be administered by other routes – such as direct
- 30 injection.

Preferably, the bacteria are killed so as to avoid the difficulties of maintaining live products.

IDENTIFYING A BACTERIUM THAT MODULATES A CELLULAR IMMUNE
5 RESPONSE

In another aspect, the present invention relates to a method for identifying one or more whole cells of bacteria from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* that modulate (e.g. modify) a cellular immune
10 response comprising the steps of: (a) contacting a first test animal with an immunostimulant; (b) contacting a second test animal with an immunostimulant mixed with a bacterium; (c) measuring the cellular immune response in each of the test animals; and (d) comparing the cellular immune response in each of the test animals, wherein, a lower cellular immune response from the immunostimulant mixed with a
15 bacterium in comparison to the immunostimulant alone is indicative of a modification of the cellular immune response by the bacterium.

In another aspect, the present invention relates to a method of determining the Th1/Th2 response of a species of bacteria selected from the genera *Rhodococcus*, *Gordonia*,
20 *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* which method comprises utilisation of the tuberculin skin test. In mice, the tuberculin skin test is preferably carried out on the foot pad. In a predominant Th1 reaction the positive foot pad immune response is maximal at 24 hours and diminishes at 48 hours. However, as the Th2 reactivity increases then the 48 hour positive foot pad immune response increases and can even exceed the
25 foot pad immune response at 24 hour.

The effect of BCG vaccination is well documented using this tuberculin skin test. Thus, the test assay can be used to assess whether or not the introduction of an immune modulator composition according to the present invention modulates the BCG cellular
30 immune response.

As used herein, the term "test animal" refers to any animal that elicits a cellular immune response to the immunostimulant. Preferably, the test animal(s) is a mammal. More preferably, the test animal(s) is a rat, hamster, rabbit, guinea pig or mouse. More preferably, the test animal(s) is a mouse.

5

Preferably, the bacterium modifies the T helper cell response. Suitably, the bacterium may modify the T helper cell response by decreasing the Th1 and Th2 response. Suitably, the bacterium may modify the T helper cell response by increasing the Th1 response and decreasing the Th2 response. Suitably, the bacterium may modify the T
10 helper cell response by increasing the Th1 response without affecting the Th2 response.

Preferably, the immunostimulant will have a known Th1 and Th2 response. For example, with the immunostimulant BCG the reaction is usually largest at 24h when it
15 is an indicator of the Th1 response; the reaction at 48h is usually less and includes a Th2 contribution. It is known that BCG predominantly stimulates a Th1 response.

By use of such immunostimulants it may be possible to determine the Th1/Th2 response of a test bacterium and, thus, it may be possible to identify one or more bacteria which have a desired Th1/Th2 response to treat and/or prevent a particular
20 disease and/or disorder.

Preferably, the cellular immune response is measured using the tuberculin skin test. Vaccination with an immunostimulant – such as BCG - induces a response to skin-testing with tuberculin (a soluble preparation of *Tubercle bacilli*), when tested later.
25 The local reaction is measured at various intervals, for example, 24 hours, 48 hours and 72 hours after injection of tuberculin. Briefly, an immunostimulant (eg. BCG) is used that induces a positive immune response to tuberculin. In the test animal, the tuberculin skin test is preferably carried out on the foot pad. In a predominant Th1 reaction the positive foot pad immune response is usually maximal at 24 hours and
30 diminishes at 48 hours. However, as the Th2 reactivity increases then the 48 hour positive foot pad immune response increases and can even exceed the foot pad immune response at 24 hour. Thus, the assay can be used to assess whether or not the

introduction of an immune modulator composition according to the present invention modulates the cellular immune response.

Preferably, the immunostimulant is BCG.

5

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph demonstrating the results of weight gain means/litter females test and controls over 12 weeks from weaning.

10

Figure 2 is a graph demonstrating the results of weight gain mean/litter males test and controls over 12 weeks from weaning.

15

Figure 3 is a graph demonstrating that *Gordonia bronchialis* enhances the early TH1 effect.

Figure 4 is a graph demonstrating that *Tsukamurella inchoensis* enhances the early TH1 response and suppresses the late TH2 response

20

Figure 5 is a graph demonstrating that *Rhodococcus coprophillus* suppresses both the TH1 and TH2 responses

Figure 6 is a graph demonstrating the tuberculin response to BCG after 1 month compared to non vaccinated controls.

25

Figure 7 is a graph demonstrating the immune modulation of the BCG effect by three species of bacteria. Tuberculin responses are measured at 24 and 48 hours. Representative species are (Na) *Nocardia asteroides*; (Gb) *Gordonia bronchialis*; and (Tp) *Tsukamurella inchoensis*.

Figure 8 is a graph demonstrating the immune modulation of BCG/BCG+ by two species of bacteria. Tuberculin responses are measured at 24, 48 and 72 hours. Representative species are (Rrh) *Rhodococcus rhodocrous* and (Dm) *Dietzia maris*.

5

Figure 9 is a graph demonstrating the immune modulation of the BCG effect by selected species within the genus *Rhodococcus*. Tuberculin responses are measured at 24, 48 and 72 hours. (Rrh) *Rhodococcus rhodocrous*; (Rru) *Rhodococcus ruber*; (Rrhod) *Rhodococcus rhodnii*; (Rcop) *Rhodococcus coprophilus*; (Ropa) *Rhodococcus opacus*; (Reryth) *Rhodococcus erythropolis*.

10

Figure 10 is a graph demonstrating the optimal dose for *Rhodococcus ruber* using BCG modified with log dilutions of *Rhodococcus ruber* from 10^8 to 10^4 compared to BCG alone.

15

The invention will now be further described by way of Examples, which are meant to serve to assist one of ordinary skill in the art in carrying out the invention and are not intended in any way to limit the scope of the invention.

20 **EXAMPLES**

METHODS

Tuberculin skin test

25

The tuberculin skin test is an appropriate model assay to assess the effect of an immune modulator composition, i.e. bacterial compositions/suspensions comprising whole killed bacterial cells according to the present invention, on a cellular immune response.

30 BCG vaccination induces a positive immune response to tuberculin. In mice, the tuberculin skin test is preferably carried out on the foot pad. In a predominant Th1 reaction the positive foot pad immune response is maximal at 24 hours and diminishes at

48 hours. However, as the Th2 reactivity increases then the 48 hour positive foot pad immune response increases and can even exceed the foot pad immune response at 24 hour.

5 The effect of BCG vaccination is well documented using this tuberculin skin test. Thus, the test assay can be used to assess whether or not the introduction of an immune modulator composition according to the present invention modulates the BCG cellular immune response.

Preparation of a bacterial suspension

10

The bacterial species from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* may be grown in an antigen-free medium, such as Sauton's medium, in a fermenter for 2-28 days. Alternatively, the bacterial species of interest may be grown on a solid slope. Alternative methods would be readily available to those skilled in the art.

15

The resulting bacterial mass may be harvested and either used directly or after washing to make a suspension in buffer. The bacterial cell suspension is prepared to contain between 100,000 and 10,000,000,000 bacilli per dose. The bacterial cells are resuspended in water or in a saline. Preferably, the saline is buffered at pH 8.0 with borate. Preferably the bacilli are inactivated (killed), suitably by heating in an autoclave for 15 minutes at 121°C. The resulting bacterial suspension comprises whole cells.

20

EXAMPLE 1: Modulation of a cellular immune response by *Rhodococcus ruber* (*R.r.*).

25

Group 1: young adult female outbred mice were left unvaccinated as a control group.

Group 2: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BCG (10^5 bacilli) (Evans).

30

Group 3: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BGC (10^5 bacilli) to which heat-killed *Mycobacterium vaccae* (M.v.) (10^7 bacilli) had been added.

- 5 All mice in Groups 1-3 were foot pad tested for tuberculin immune response at days 10 and 30. Then each mouse was injected with heat-killed *Mycobacterium vaccae* (10^7 bacilli) at day 40. At day 50 the tuberculin test on the foot pad was repeated.

Group 4: young adult female outbred mice were left unvaccinated as a further control group.

Group 5: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BCG (Evans) (10^5 bacilli).

Group 6: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BGC (10^5 bacilli) to which heat-killed *Mycobacterium vaccae* (10^7 bacilli) had been added.

Group 7: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BGC (10^5 bacilli) to which heat-killed *Rhodococcus ruber* (10^7 bacilli) had been added.

Group 8: young adult female outbred mice were vaccinated on day 0 into the scruff of the neck with BCG (10^5 bacilli) to which heat-killed *Rhodococcus ruber* (10^6 bacilli) had been added.

All mice in Groups 4-7 were foot pad tested for tuberculin immune response at day 30. Then at day 40 each group was divided so that half of each group received no further treatment and the second half of each group received an injection of *Rhodococcus ruber* (10^7 bacilli). At day 50 the tuberculin test on the foot pad was repeated.

The result of Example 1 are shown in Table 1.

TABLE 1: Modulation of a cellular immune response by *Rhodococcus ruber* (R.r.)

		Tuberculin response at		
Day		24 hours	48 hours	24-48 difference
5	Group 1 - control			
	10	6.9±4.5	5.3±3.7	1.6
	30	5.9±4.7	5.2±4.5	0.7
	50 M.v.	13.1±5.6 p<0.01	14.0±5.2 p<0.001	+0.9
10	Group 2 - BCG			
	10	16.8±11.1	16.8±5.5	0
	30	31.1±17.8	16.9±9.1	14.2 <0.05
	50 M.v.	33.9±12.3	19.2±11.3	14.7 <0.02
15	Group 3- BCG + M.v.			
	10	14.6±8.3	9.9±2.7	4.7
	30	23.8±16.1	21.7±11.5	2.1 n.s.
	50 M.v.	27.4±10.3	15.1±5.8	12.3 <0.005
20	Group 4 - control			
	30	2.8±3.7	2.5±2.3	0.3 n.s.
	50	6.4±9.4	3.6±7.5	2.8 n.s.
	50 R.r.	4.2±5.4	1.0±1.7	3.2 n.s.
25	Group 5- BCG			
	30	28.8±15.4	19.1±10.9	9.7 (<0.1)
	50	33.8±21.6	12.0±7.9	21.8 (<0.1)
	50 R.r.	40.0±22.4	12.2±9.2	27.8 <0.05
30	Group 6- BCG + M.v.			
	30	19.4±20.5	17.7±11.8	1.7 n.s.
	50	56.8±53.8	29.4±31.1	27.4 n.s.
	50 R.r.	50.0±38.4	9.0±10.4	41.0 <0.05
35	Group 7- BCG + R.r.			
	30	20.6±10.9	17.8±10.3	2.8 n.s.
	50	24.8±22.5	14.6±13.2	10.2 n.s.
	50 R.r.	28.0±13.2	11.0±4.7	17.0 <0.05
40	Group 8 - BCG + R.r./10 group			
	30	19.5±13.9	20.9±14.9	+1.4
	50	37.2±25.2	20.2±5.3	17.0 n.s.
	R.r.			
45	50	26.8±14.8	12.8±8.0	14.0 (<0.1)
	M.v. = <i>Mycobacterium vaccae</i>			

In the control Group 1 *Mycobacterium vaccae* treatment induced a statistically significant increase in the immune response to tuberculin after both 24 hours ($p < 0.01$) and 48 hours ($p < 0.001$). However, in the control Group 4, treatment with *Rhodococcus ruber* did not induce a significant change in immune response to tuberculin after both 24 hours and 48 hours. At both time points the *M.v.* results were significantly greater than the *R.r.* results ($p < 0.02$).

In the BCG groups (Group 2 and Group 5), the fall in response to tuberculin between 24 hours and 48 hours was greater ($p = 0.06$) in the mice receiving treatment with *Rhodococcus ruber* (the mean fall being 28.2 ± 15.7) than in the mice receiving treatment with *Mycobacterium vaccae* (the mean fall being 14.9 ± 9.6).

In the BCG+*Mycobacterium vaccae* group (Group 3 and Group 6), the fall in response to tuberculin between 24 hours and 48 hours was again greater ($p < 0.05$) in the group receiving treatment with *Rhodococcus ruber* (mean fall being 41.0 ± 41.0) than in mice receiving treatment with *Mycobacterium vaccae* (mean fall being 12.7 ± 7.0).

These data suggest that there is a down-regulation of the Th2 response after treatment with *Rhodococcus ruber*, which is not seen following treatment with *Mycobacterium vaccae*.

In the BCG + *R.r.* groups (Groups 7 and 8) the effect of adding *R.r.* 10^7 (Group 7) or 10^6 (Group 8) to BCG was very similar and after the second injection with *R.r.* there was a substantial reduction in response between 24 and 48 hours (15.5 ± 10.0).

EXAMPLE 2: Modulation of a cellular immune response using *Nocardia asteroides* (*N.a.*), *Gordonia bronchialis* (*G.b.*) or *Tsukamurella inchoensis* (*T.p.*).

This experiment was designed to compare the effects of foot-pad testing with Tuberculin 28 days after vaccinating groups of 6 mice with BCG alone or with the addition of 10^7 *M.v.*, *R.r.*, *Nocardia asteroides* (*N.a.*), *Gordonia bronchialis* (*G.b.*) or *Tsukamurella inchoensis* (*T.p.*). To find out what the effects were on the immune response to each of

these added organisms, the groups of animals were tested in the other foot-pad 28 days after they had been tested with Tuberculin with skin-test reagents made from the organism included in their vaccination (Vaccin, Rubin, Asterin, Bronchialin and Inehonensin).

5 The results are detailed in Table 2.

TABLE 2: Modulation of a cellular immune response

		Tuberculin response at		
		24 hours	48 hours	24-48 h difference
10	Day	1) Control group		
	28	3.8±4.8	3.7±3.6	0.1
15	28	2) BCG group		
	Tuberculin	38.0±20.2	28.2±17.0	9.8
	56	65.0±31.2	45.8±23.6	19.2
20	28	3) BCG+<i>M.v.</i> group		
	Vaccin	20.3±10.0	14.2±5.7	6.1
	56	18.7±12.1	8.0±6.5	10.7
25	28	4) BCG+<i>R.r.</i> group		
	Rubin	31.3±16.0	24.2±10.3	7.1
	56	19.5±7.1	12.3±10.4	7.2
30	28	5) BCG+<i>N.a.</i> group		
	Asterin	24.2±20.8	20.2±17.6	4.0
	56	7.3±8.3	5.3±5.0	2.0
35	28	6) BCG+<i>G.b.</i> group		
	Bronchialin	15.8±14.4	15.7±13.2	0.1
	56	11.3±3.4	6.2±4.3	5.1
40	28	7) BCG+<i>T.p.</i> group		
	Paurometabolin	19.5±7.4	15.8±5.7	3.7
	56	9.8±5.0	2.3±2.7	7.5

All of the bacterial suspensions depressed the 28 day Tuberculin response measured at both 24 and 48 hours, in comparison with that following BCG alone ($p=0.05$; $p=0.2$).

- 5 With the exception of Tuberculin and Vaccin, this was the first time that any of these skin test reagents have been used. The differences in responsiveness to the new reagents at 24 hours were probably because they had not been equilibrated except by protein estimation. However, all showed a fall in response between 24 and 48 hours at
10 day 50, suggesting immunoregulatory activity.

EXAMPLE 3: Modulation of a cellular immune response by *Rhodococcus ruber* treatment

- 15 This experiment was designed to investigate the effect of Diphtheria/Tetanus/Pertussis (DTP) vaccine and/or Measles/Mumps/Rubella (MMR) vaccine on subsequent BCG vaccination and how the effect can be modified by treatment with heat killed whole cells of *Rhodococcus ruber* given 7 days or 28 days in advance, or with the first dose of DTP.
- 20 The experimental design is detailed in Table 4. Groups of six weaning female outbred mice received the treatments detailed in Table 4. All mice were tuberculin-tested 28 days after BCG vaccination.

The results are presented in Table 3.

TABLE 3:

		24 hours	48 hours	24-48 h difference
	Group 5 in Table 4			
5	BCG only:	35.0 \pm 24.7	22.0 \pm 14.8	13.0
	Group 3 in Table 4			
	DTP/MMR			
	then BCG:	42.7 \pm 28.5	24.2 \pm 13.2	18.5
10				
	Group 6 in Table 4			
	R.r. given 49 days			
	before BCG:	50.3 \pm 47.0	31.8 \pm 32.0	18.5
15				
	Group 1 in Table 4			
	R.r. given 7 days			
	before DTP/MMR:	32.0 \pm 18.4	24.3 \pm 15.4	7.7
	Group 2 in Table 4			
20	R.r. given 28 days			
	before DTP/MMR:	44.7 \pm 25.8	20.0 \pm 11.0	24.7
	Group 4 in Table 4			
	DTP/MMR			
25	then BCG:	42.0 \pm 21.1	23.7 \pm 8.0	18.3
	Group 7 in Table 4			
	R.r. DTP/MMR			
	then BCG:	40.3 \pm 26.0	27.3 \pm 16.5	13.0
30				

Trends shown in this experiment:

The effect of giving DTP/MMR before BCG (Groups 3 & 4 in Table 4): the administration of DTP and MMR before vaccination with BCG results in an increase in the 24 hour response and better separation between the 24 and 48 hour responses.

5

The effect of giving *R.r.* before BCG (Group 6 in Table 4): *R.r.* given 49 days before BCG increases the Tuberculin response at 24 and 48 hours, and increases the difference between them.

10 The effect of giving *R.r.* before DTP/MMR before BCG (Groups 1 & 2 in Table 4): giving *R.r.* 28 days before DTP/MMR increases the separation between the 24 and 48 hour Tuberculin responses, whereas giving *R.r.* 7 days before DTP/MMR suppresses the effect on subsequent BCG.

15 The effect of giving *R.r.* with the first dose of DTP/MMR before BCG (Group 7 in Table 4): giving *R.r.* with the first injection of DTP appeared to make no difference to the effect of DTP/MMR on subsequent BCG vaccination.

20 Preliminary findings suggest that immunisation with DTP followed by MMR, prior to BCG vaccination, has an effect on Tuberculin reactivity 28 days later. The study also suggests that vaccination with *R.r.* prior to BCG vaccination, has an effect on Tuberculin reactivity 28 days later. The timing of vaccination with *R.r.* in relation to DTP/MMR prior to BCG vaccination may be important.

25 **EXAMPLE 4: Local skin reactions to intradermal injections in adult Guinea Pigs**

The left flank of 3 animals was shaved to give intradermal injections of 0.1 ml containing 10^8 *M. vaccae* at the head end and 0.1ml containing 10^9 *M. vaccae* 5cm towards the tail end of the animal.

30

Another 3 animals were shaved in the right flank and given intradermal injections of 0.1 ml containing 10^8 *R. rhodochrous* at the head end and 0.1 ml containing 10^9 *R. rhodochrous* 5cm-towards the-tail end-of-the animal.

5 Diameters of induration in mm

Groups	48 h 10^8	7d 10^8	14d 10^8	48 h 10^9	7d 10^9	14d 10^9
<i>M. vaccae</i>	2x2	-	-	4x4	2x3	-
<i>M. vaccae</i>	-	-	-	2x2	2x2	1x1
<i>M. vaccae</i>	-	-	-	5x5	1x2	-
<i>R. rhodochrous</i>	-	-	-	1x1	2x2	-
<i>R. rhodochrous</i>	-	-	-	1x1	2x2	-
<i>R. rhodochrous</i>	-	-	-	2x1	4x3	1x2

In 3 guinea pigs, the local reactions to intradermal injections (the route of injection to be typically used in veterinary and medical practice) of 10^9 *R. ruber* (a typical dose for human and animal use) were similar to reactions to the same dose of *M. vaccae* in another 3 guinea pigs. At 48 hours after injection, reactions to *R. ruber* were smaller ($p < 0.05$) than those to *M. vaccae*. Thus, *R. ruber* may be even more pharmaceutically acceptable than *M. vaccae*. There are no local reactions to either preparation at the 10^8 dose.

15

EXAMPLE 5: Toxicity following subcutaneous injection.

No evidence of toxicity to subcutaneous doses was observed in 17 rats receiving 3 injections of *R. ruber* when they were aged 1 day, 14 days and 28 days (the last of these were given 7 days after challenge with *Trypanosoma cruzi* in expt. 6).

20

The responses to 10^9 *R. ruber* at 48 hours were smaller than those to the same dose of *M. vaccae* ($p < 0.05$). There were no differences at 7 days or 14 days. Many mice have

received injections of various species of the bacteria of the present invention, without any evidence of toxicity.

EXAMPLE 6: Modulation of Trypanosomiasis in rats

5

Preparation of animals

- i) One day old male "1" rats were injected into the scruff of the neck with 10^7 *Rhodococcus ruber* in a volume of 0.1 ml via subcutaneous injection.
- ii) After 14 days old, the male "1" rats were given a second subcutaneous injection with 10^7 *R. ruber* in 0.1ml on the left side.
- 10 iii) Animals were challenged with live *Trypanosoma cruzi* on day 21 via the subcutaneous route with 10^6 trypomastigotes of the Tulahuén strain of *T. cruzi*. Infective blood trypomastigotes were maintained by serial passage in CBI mice.
- 15 iv) Bloodstream forms of *T. cruzi* were assessed under standardised conditions, by direct microscopic observation of 5 μ l of heparinized tail venous blood, at 7 and 14 days post-infection (pi). Data are expressed as number of parasites/50 fields.
- v) 7 days later an additional subcutaneous injection of 10^7 *R. ruber* was
20 administered in a volume of 0.1ml on the right side.

Control animals only received the challenge with *T. cruzi*.

Another group of animals was left unchallenged for comparison purposes.

25

The results are presented in Table 5.

TABLE 5: Parasitemias (Data are expressed as number of parasites/50 fields)**1st Experiment**

5	GROUP	n	Day 7		Day 14	
			Mean±SD	Median (rank)	Mean±SD	Median (R)
	Controls	7	2.57±1.8	2(1-5)	1.14±1.07	1(0-3)
	Rr	10	0.30±0.67	0(0-2)	0±0	0(0-0)
10			P<0.005 Student's t test		P<0.005	

2nd Experiment

15	GROUP	n	Day 7		Day 14	
			Mean±SD	Median (rank)	Mean±SD	Median (R)
	Controls	7	2.14±1.07	2(1-4)	1±1.15	1(0-3)
	Rr	7	1.1±0.69	0(1-2)	1.5±0.97	2(0-3)
20			P=0.05 Student's t test			
25						

Pooled data from both experiments

30	GROUP	n	Day 7		Day 14	
			Mean±SD	Median (rank)	Mean±SD	Median (R)
	Controls	14	2.35±1.44	2(1-5)	1.07±1.07	1(0-3)
	Rr	17	0.65±0.78	0(1-2)	0.69±1.01	0(0-3)
35			P=0.001 Student's t test			

This data, demonstrates a significant reduction of parasites circulating in the blood at the time when the greatest numbers are found in the control animals. This suggests that *R. ruber* enhances the Th1 responses (7 days is too early for antibody production via the Th2 route).

5

EXAMPLE 7: Investigating the effects of *Rhodococcus coprophilus* NCIMB 11211, *Gordonia bronchialis* NC10667 and *Tsukamurella inchoensis* NC13040 on toxicity, in particular growth rate and subsequent BCG vaccination in newborn mice.

10

An alternative test of toxicity is that in which subcutaneous injections of *Gordonia bronchialis*, *Rhodococcus coprophillus* or *Tsukamurella inchoensis* are given on the day of birth, and again in a separate site, on day 21. These animals, and saline-injected controls were weighed at regular intervals over 3 months.

15

Methods

9 late term pregnant Balb C mice are used.

20 Preparation of killed bacterial suspension: Suspension prepared from 10 day Sauton antifoam broth culture, centrifuged and resuspended as a 10 mgs/ml in borate buffered saline, autoclaved and stored at 4°C. Dilute 10 mgs/ml, 10^{10} /ml 1/2 in borate giving 5×10^9 and then diluting 1/10 in borate buffer giving a final concentration of 10^7 in 20 μ for use.

25

Day 1

On day of birth remove newborn mice singly and inject litter with 10^7 killed suspension into the scruff and return to mother.

30

Inject control litters with 20 μ litres of M15 borate buffered saline into the scruff.

Litter 1 = 2 female + 3 male (*Rhodococcus coprophilus* NCIMB 11211)

Litter 2 = 2 female + 3 male (*Rhodococcus coprophilus* NCIMB 11211)

5 **Litter 3** = 5 female + 3 male (*Gordonia bronchialis* NC10667)

Litter 4 = 3 female + 4 male (*Gordonia bronchialis* NC10667)

Litter 5 = 1 female + 1 male (Borate buffered saline)

10 **Litter 6** = 3 female + 1 male (Borate buffered saline)

Litter 7 = 5 female + 3 male (*Tsukamurella inchoensis* NC13040)

Litter 8 = 5 female + 4 male (*Tsukamurella inchoensis* NC13040)

15

Litter 9 = 4 female + 5 male (borate buffered saline)

Day 21

20 Wean the mice, sex and separate the litters into 2 groups male and female

Weigh and revaccinate. Mark individuals by tail marking with an indelible pen.

Weigh and re tail mark weekly for 3 months.

25

Figure 1 demonstrates the results of weight gain means/litter females test and controls over 12 weeks from weaning.

Figure 2 demonstrates the results of weight gain mean/litter males test and controls over 12 weeks from weaning.

30

Weight gain is the same in all groups, and only 2/57 animals died over the 3 months. Both were males, that died just after being put together with other males after weaning. Thus it can be concluded that weight gain and progress of the animals was unaffected by the three species tested.

5

EXAMPLE 8: The effect on BCG immunisation measured through the tuberculin skin test response

10 An immune modulator test model is devised, based upon on the principle that vaccination with BCG induces a response to skin-testing with Tuberculin (a soluble preparation of *Tubercle bacilli*), when tested 4 weeks later. The local reaction is measured 24 hours, 48 hours and 72 hours after injection of Tuberculin. In the mouse the reaction is usually largest at 24 hours when it is an indicator of the Th1 response to the antigens in Tuberculin. The reaction at 48 hours is usually less and includes a Th2
15 contribution. The reaction at 72 hours is often little less than at 48 hours and is a Th2 response. This post-BCG Tuberculin reaction can be modulated by prior priming, so that the Th1 and Th2 components of the reaction will reflect the nature of the priming reagent.

20 Methods

Mice primed according to Example 8 when 3 months old were injected with 10^6 BCG in 100 μ l into the scruff to half of each test and the control group.

25 At 4 months tuberculin test with reagent T1475 1mg/ml. Dilute 100 μ litres in 1.9 ml to give final concentration of 50 μ gms/ml. Store at 4 °C. Dose is 2.5 μ g in 50 μ litres given into the hind footpad. Tuberculin response swelling as measured using a micrometer at 24, 48 and 72 hours.

Results

Figure 3 shows that *Gordonia bronchialis* enhances the 24 hour TH1 effect.

Figure 4 shows that *Tsukamurella inchoensis* enhances the 24 hour TH1 response and suppresses the 72 hour TH2 response

5

Figure 5 shows that *Rhodococcus coprophillus* suppresses both the TH1 and TH2 responses

Conclusion

10

The 3 species each induced different effects on the tuberculin test following BCG challenge:

Gordonia bronchialis induced an enhanced Th1 response, without changing the Th2 response. *Rhodococcus ruber* also has this function.

15

Tsukamurella inchoensis enhanced the Th1 response and down-regulated the Th2 response.

Rhodococcus coprophilus down-regulated both Th1 and Th2 responses.

25

EXAMPLE 9: Immune modulator test model

An immune modulator test model is devised, based upon on the principle that vaccination with BCG induces a response to skin-testing with Tuberculin (a soluble preparation of *Tubercle bacilli*), when tested 4 weeks later. The local reaction is measured 24 hours, 48 hours and 72 hours after injection of Tuberculin. The reaction is usually largest at 24 hours when it is an indicator of the Th1 response to the antigens

30

in Tuberculin. The reaction at 48 hours is usually less and includes a Th2 contribution. The reaction at 72 hours is often little less than at 48 hours and is a Th2 response. This post-BCG Tuberculin reaction can be modulated by prior priming, so that the Th1 and Th2 components of the reaction will reflect the nature of the priming reagent.

5

The known immunostimulant, BCG is injected into the scruff of young 3 week old mice and the tuberculin response measured 1 month later by subcutaneous injection of tuberculin into the mouse footpad. The resultant swelling ie. the "tuberculin response" is then measured at 24, 48 and 72 hours. Swelling at 24 hours is considered an early or Th1 mediated response and swelling at 48 and 72 hours a late or Th2 mediated response. BCG in the healthy mouse stimulates predominantly a Th1 response.

10

Method

(a) BCG intradermal vaccine 10 dose vial (Evans Medical).

15 Reconstitute with 1 ml supplied sterile water using a syringe and needle allowing 5 minutes to dissolve. Should be 1×10^7 /ml.

Using a syringe and needle remove all of the vaccine and transfer to a plastic bijou bottle.

20

Dilute 1/10 0.15ml in 1.35ml M15 borate buffered saline gives 10^5 in 100µlitres

Dose is 10^5 in 100 µl given into the scruff of the neck.

25 (b) Tuberculin

T1475 1mg/ml.

Dilute 100µl in 1.9 ml to give final concentration of 50µgms/ml.

Store at 4 °C.

Dose is 2.5µg in 50µl given intradermally into the hind footpad.

5

Tuberculin response is measured at 24, 48 and 72 hours using a micrometer

Figure 6 shows the tuberculin response to BCG after 1 month compared to non vaccinated controls.

10 (c) Preparation of Test Suspensions

Cultures are grown in Sauton broth harvested by centrifugation and resuspended at a concentration of 10mg/ml in M15 borate buffered saline and stored at 4°C.

15 10mgs/ml = 10^9 in 100µl.

Dilute 1/10 = 10^8 in 100µl.

Add 150µl 10^8 to 1.2ml M15 borate and add 150µl BCG.

20

Dose is now 10^5 BCG + 10^7 test organism in 100µl injected into the scruff of the neck.

Results

25 Figure 7 is a graph demonstrating the immune modulation of the BCG effect by three species of bacteria. Tuberculin responses are measured at 24 and 48 hours. Representative species are (Na) *Nocardia asteroides*; (Gb) *Gordonia bronchialis*; and (Tp) *Tsukamurella inchonensis*.

Figure 8 is a graph demonstrating the immune modulation of BCG/BCG+ by two species of bacteria. Tuberculin responses are measured at 24, 48 and 72 hours. Representative species are (Rrh) *Rhodococcus rhodocrous* and (Dm) *Dietzia maris*.

5

Figure 9 is a graph demonstrating the immune modulation of the BCG effect by selected species within the genus *Rhodococcus*. Tuberculin responses are measured at 24, 48 and 72 hours. (Rrh) *Rhodococcus rhodocrous*; (Rru) *Rhodococcus ruber*; (Rrhod) *Rhodococcus rhodnii*; (Rcop) *Rhodococcus coprophilus*; (Ropa) *Rhodococcus opacus*; (Reryth) *Rhodococcus erythopolis*.

10

Figure 10 is a graph demonstrating the optimal dose for *Rhodococcus ruber* using BCG modified with log dilutions of *Rhodococcus ruber* from 10^8 to 10^4 compared to BCG alone.

15

Conclusion

This screening test demonstrates that by mixing test suspensions with BCG and comparing to BCG alone, 28 days later the tuberculin test, as a measure of response to BCG, has been regulated or modified. In this simple model, immunoregulation is shown as a depression in the size of the response at both 24 and 48 hours. This can be further investigated in longer term experiments in which priming with the test suspension is carried out some weeks before challenge with BCG and subsequent Tuberculin testing.

25

EXAMPLE 10: Growth enhancement in pigs.

This is based on the principle that an immune system primed for Th1 enhancement, with or without regulation of Th2 results in less restriction of growth by subclinical infections. Thus in a situation where minor infections are frequent, a reduction of their influence should result in the animals putting on more weight. This might be achieved by reducing the level of circulating TNF (Tumour Necrosis Factor), which in the

30

presence of Th2 cytokines acts as "cachectic factor", reducing appetite and increasing the metabolic rate.

Two litters each of a pig strain producing 11-15 piglets per litter will receive 3
 5 intradermal injections of 0.1ml of test reagent or placebo, at the ages of 0 (day of birth),
 aged 3 weeks and 64 days when they are mixed with other piglets. Animals will be
 individually weighed at 3 weeks, at 64 days and at marketing ages 4-5 months.

Preliminary investigations suggest that piglets receiving the different test preparations
 10 show varying degrees of weight enhancement in comparison with the control group
 given similar injections, but of buffered saline.

EXAMPLE 11: Modulation of Sweet Itch in horses

15 Sweet-itch is the skin condition commonly troubling horses and ponies making an
 allergic response to bites by *Culicoides* midges. This most commonly occurs in areas
 where hair is long, such as the mane and tail. There is a loss of hair and thickening and
 inflammation of the skin with intense itching. Horses are excited and distressed by this
 condition and may lose weight as a result thereof. This is an annual condition affecting
 20 afflicted horses and ponies in the late spring and early summer usually.

Groups of approximately 5 to 15 ponies and horses known to develop sweet-itch every
 year, will receive *T. inchoensis*, *R. coprophilus*, a positive control or a placebo
 control. Two injections of each will be given to each animal, the first in late February
 25 and the other three weeks later. Observations of behaviour, hair condition and skin
 condition will be made for the following three months at monthly intervals, and again
 at the end of the midge season if this is prolonged. Quantitative and qualitative
 measurements will be developed to assess progress. Observers will be blind as to
 which treatment each group of animals has received.

Preliminary investigations suggest that at least one and possibly both test organisms will result in measurably reduced hair loss and skin inflammation in comparison with the placebo control.

5 **EXAMPLE 12: Cancer model**

This study assesses the effect of injections of 10^6 and 10^7 *Gordonia bronchialis*, *Rhodococcus coprophilus* and *Tsukamurella inchoensis* in the prevention and treatment of mice challenged with a lung adenocarcinoma cell line. Groups of mice will receive subcutaneous injections of the test species or saline placebo into the scruff of the neck 5 weeks and two weeks before subcutaneous challenge with tumour cells into the side. The third injection of the test organisms and saline placebo will be given into the scruff of the neck 1 week after tumour challenge. The number of tumour cells injected will be decided by those fully conversant with the system. End points will be the presence and size of the tumours at the challenge site 7, 14 and 21 days after challenge.

Preliminary investigations suggest that at least one of the test bacteria will delay tumour development and reduce tumour progression.

20 **Example 13: Comparison of the cellular immune response obtained using cell sonicates as compared with whole cells of the genera of the present invention.**

To show that the suspensions of whole cells of the bacteria are more effective in modulating immunity than cell extracts, comparative preparations will be made and tested in a BCG challenge/Tuberculin skin test system.

60 mg of *Rhodococcus coprophilus* will be suspended in buffered saline (pH8.0) at a concentration of 10 mg wet weight of organisms/ml. After autoclaving at 121 °C for 15 minutes, 3 ml will be treated in an ultrasonicator to break open the majority of organisms (70-80%).

A measured volume of the sonicate will be centrifuged at 4000 rpm for 15 mins in a pre-weighed tube and the supernatant will be carefully removed as the test extract. The deposit will be spun again and the remaining supernatant drained off. The tube will be weighed to determine the proportion of whole organisms that were not broken open. The
 5 volume of extract equivalent to 10^7 *R. coprophilus* will then be compared with the equivalent whole bacilli.

Groups of 10 animals will receive injections of the extract of bacilli or whole cell suspensions (10^7 /dose) or saline placebo into the scruff of the neck on the day of birth and
 10 again aged 21 days. 8-12 weeks later animals will be challenged with BCG. 28 days later Tuberculin testing will be performed with readings taken at 24, 48 and 72 hours.

Preliminary investigations suggest that whole cells of the bacteria are more effective in modulating immunity than cell extracts.

15

Example 14: Stress in Pigs

To evaluate the effect of whole cells of bacteria from one or more of the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioides* on stress
 20 in pigs (piglets), one or more species from one or more of these genera were tested in pigs.

Two litters each of a pig strain producing 11-15 piglets per litter will receive 3 intradermal injections of 0.1ml of test reagent or placebo, at the ages of 0 (day of
 25 birth), aged 3 weeks and 64 days when they are mixed with other piglets.

One or more saliva swabs will be taken from each animal at 3 weeks, at 64 days and at marketing aged 4-5 months. The saliva will be analysed for stress-related hormones, in particular cortisol.

30

It is expected that one or more of the test bacteria will reduce stress by neuro-endocrino-immunological mechanisms.

All publications mentioned in the above specification are herein incorporated by reference. Various modifications and variations of the described methods and system of the present invention will be apparent to those skilled in the art without departing from the scope and
5 spirit of the present invention. Although the present invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention which are obvious to those skilled in biochemistry and biotechnology or related fields are intended to
10 be within the scope of the following claims.

Table 4: Experimental Design. Groups of 6 weanling female outbred mice received the following treatments:

	Day 0	day 7	day 14	Day 21	Day 28	day 35	day 42	day 49	day 56	day 77	Day 84	Day 105
Group 1	R/r	DTP	DTP	DTP	MMR				BCG		TT	
Group 2	R/r				DTP	DTP	DTP	MMR		BCG		TT
Group 3		DTP	DTP	DTP	MMR				BCG		TT	
Group 4					DTP	DTP	DTP	MMR		BCG		TT
Group 5									BCG		TT	
Group 6					R/r					BCG		TT
Group 7		R/r/DTP	DTP	DTP	MMR				BCG		TT	

Key

R/r Rhodococcus ruber
 DTP Diphtheria/tetanus/pertussus
 MMR Measles/Mumps/Rubella
 BCG BCG vaccine
 TT Tuberculin Test

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CLAIMS

1. An immune modulator composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea*, wherein said immune modulator composition in use modifies a cellular
5 immune response.
2. An immune modulator composition according to claim 1 wherein said composition comprises an antigen and an adjuvant, wherein said adjuvant comprises a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*,
10 *Tsukamurella* and *Nocardioidea*.
3. A pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*, *Tsukamurella* and *Nocardioidea* and optionally a pharmaceutically acceptable carrier, diluent or excipient, which
15 pharmaceutical composition in use modifies a cellular immune response.
4. Use of an immune modulator composition or a pharmaceutical composition according to any one of claims 1 to 3 in the manufacture of a medicament for the treatment or prevention of one or more of an infection (e.g. a bacterial, viral, or
20 parasitic infection, for example, malaria, trypanosomiasis, leishmaniasis and toxoplasmosis) and/or the immunological abnormalities accompanying an infection, stress, for example, major trauma stress, psychosocial stress and chronic stress, an allergy (e.g. asthma including allergic asthma, hayfever, allergic dermatitis (eczema), anaphylactic shock, allergies to plant contact or ingestion, stings – such as nettle and
25 insect stings, and allergies to insect bites – such as midges, for instance *Culicoides* (which causes Sweet Itch in horses)), heaves, COPD and cancer (for example melanoma or adenocarcinoma) and an immune system imbalance (e.g. an immune system imbalance in children and the elderly).
5. Use of an immune modulator composition or a pharmaceutical composition
30 according to any one of claims 1 to 3 in the manufacture of a medicament for the

treatment or prevention of one or more of an infection, stress, an allergy, heaves, COPD, cancer and an immune system imbalance.

6. Use of an immune modulator composition or a pharmaceutical composition
5 according to any one of claims 1 to 3 in the manufacture of a medicament for the
treatment or prevention of an adverse reaction to childhood vaccines and/or
consequences thereof.

7. Use of an immune modulator composition or a pharmaceutical composition
10 according to any one of claims 1 to 3 in the manufacture of a medicament for
modifying a cellular immune response.

8. An immune modulator composition or a pharmaceutical composition
according to any one of claims 1 to 3 for use as a medicament.

15

9. An immune modulator composition or a pharmaceutical composition
according to any one of claims 1 to 3 for use in or as a vaccine.

10. An immune modulator composition according to claim 9 wherein said
20 vaccine is a prophylactic vaccine or a therapeutic vaccine.

11. An immune modulator composition according to claim 9 or claim 10
wherein said composition modifies a cellular immune response.

25 12. An immune modulator composition and/or a pharmaceutical composition
according to any one of the preceding claims wherein said composition further
comprises an antigen or antigenic determinant.

30 13. An immune modulator composition and/or a pharmaceutical composition
according to claim 12 wherein said antigen or antigenic determinant is an antigen or
antigenic determinant selected from one or more of a BCG (bacillus of Calmette and

Guerin) vaccine, a diphtheria toxoid vaccine, a diphtheria/tetanus/pertussis vaccine, a pertussis vaccine, the tetanus toxoid vaccine, the measles vaccine, the mumps vaccine, the rubella vaccine, the OPV (oral poliomyelitis vaccine) and *Mycobacterium vaccae*, or part thereof.

5

14. An immune modulator composition and/or pharmaceutical composition according to claim 12 or claim 13 wherein said composition comprises two or more such antigens or antigenic determinants.

10

15. An immune modulator composition and/or pharmaceutical composition according to any one of claims 1-3 and 8-14 wherein said bacterium is selected from the genus *Rhodococcus*.

15

16. An immune modulator composition and/or pharmaceutical composition according to claim 15 wherein said bacterium is *Rhodococcus ruber*.

17. An immune modulator composition and/or pharmaceutical composition according to any one of claims 1-3 and 8-16 wherein said bacterium is killed.

20

18. A method for treating or preventing a condition in a subject comprising administering an effective amount of a pharmaceutical composition and/or immune modulator composition according to any one of claims 1-3 and 8-17 to a subject wherein said composition modulates a cellular immune response.

25

19. A method for immunising a subject comprising administering a pharmaceutical composition and/or immune modulator composition according to any one of claim 1-3 and 8-17.

30

20. A method according to claim 18 or claim 19 wherein said pharmaceutical composition and/or an immune modulator composition is co-administered with an antigen or antigenic determinant.

21. A method according to claim 20 wherein the antigen or antigenic determinant is an antigen or antigenic determinant selected from one or more of a BCG (bacillus of Calmette and Guerin) vaccine, a diphtheria toxoid vaccine, a
5 diphtheria/tetanus/pertussis vaccine, a pertussis vaccine, a tetanus toxoid vaccine, a measles vaccine, a mumps vaccine, a rubella vaccine, a OPV (oral poliomyelitis vaccine) and *Mycobacterium vaccae*, or part thereof.

22. A method according to claim 20 or claim 21 wherein said pharmaceutical
10 composition and/or an immune modulator composition is co-administered with two or more such antigens or antigenic determinants.

ABSTRACT

An immune modulator composition and/or pharmaceutical composition comprising a whole cell of a bacterium from the genera *Rhodococcus*, *Gordonia*, *Nocardia*, *Dietzia*,
5 *Tsukamurella* and *Nocardioides*, wherein said immune modulator composition in use modifies a cellular immune response.

Figure 1

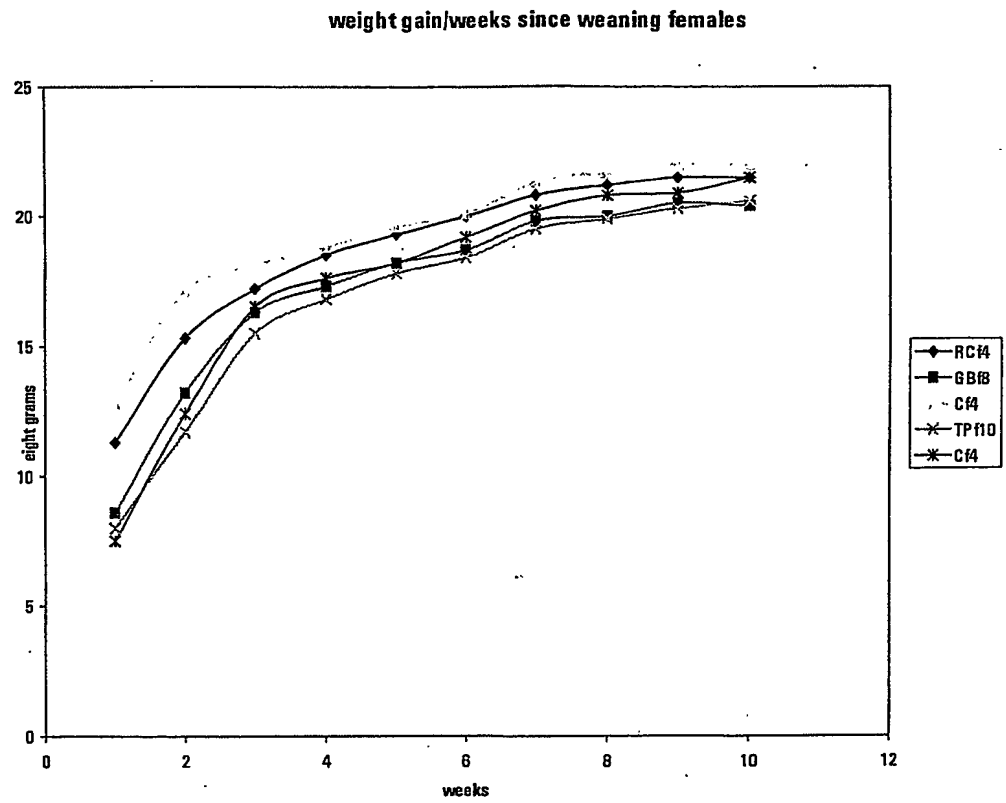


Figure 2

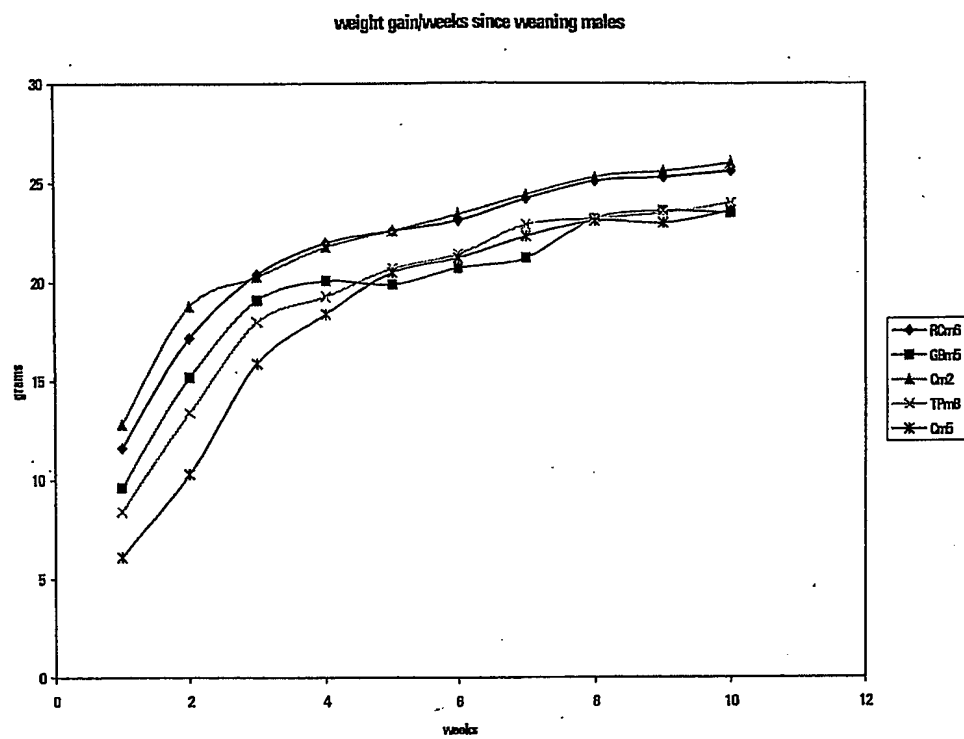


Figure 3

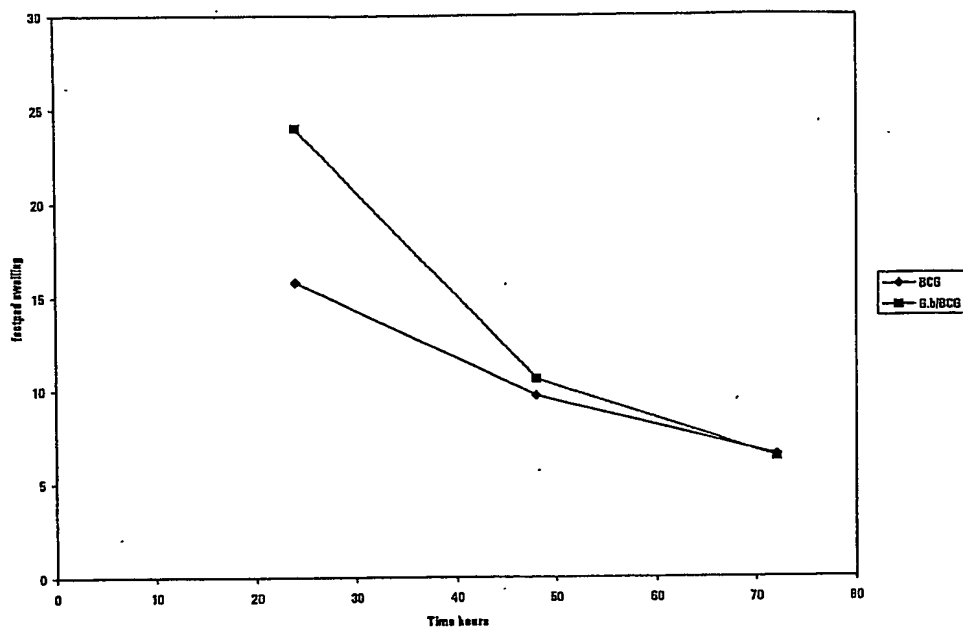


Figure 4

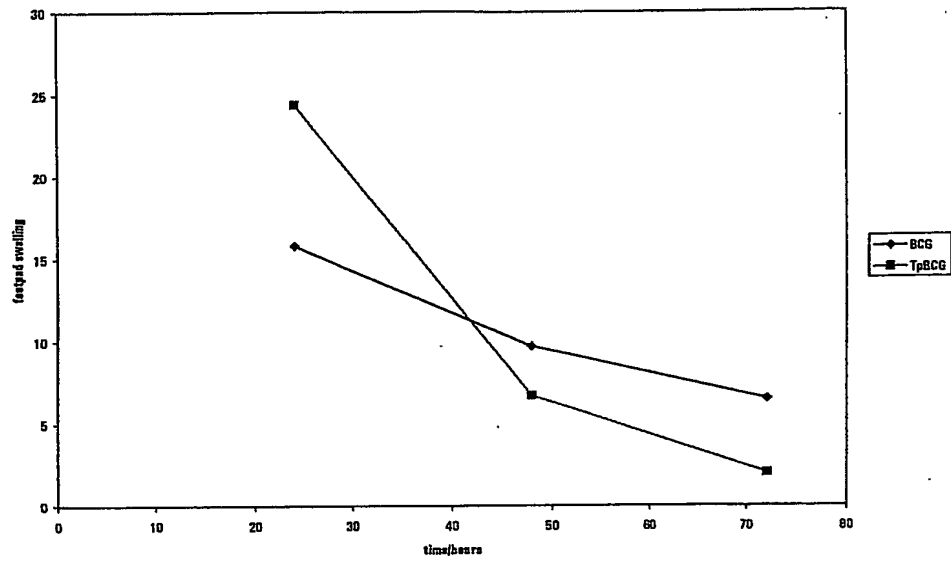


Figure 5

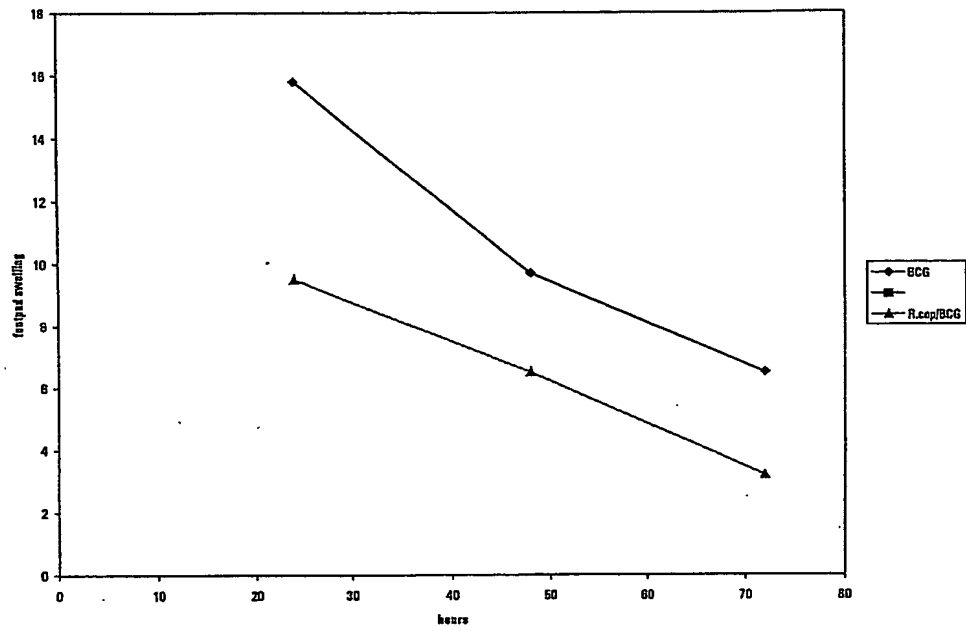


Figure 6

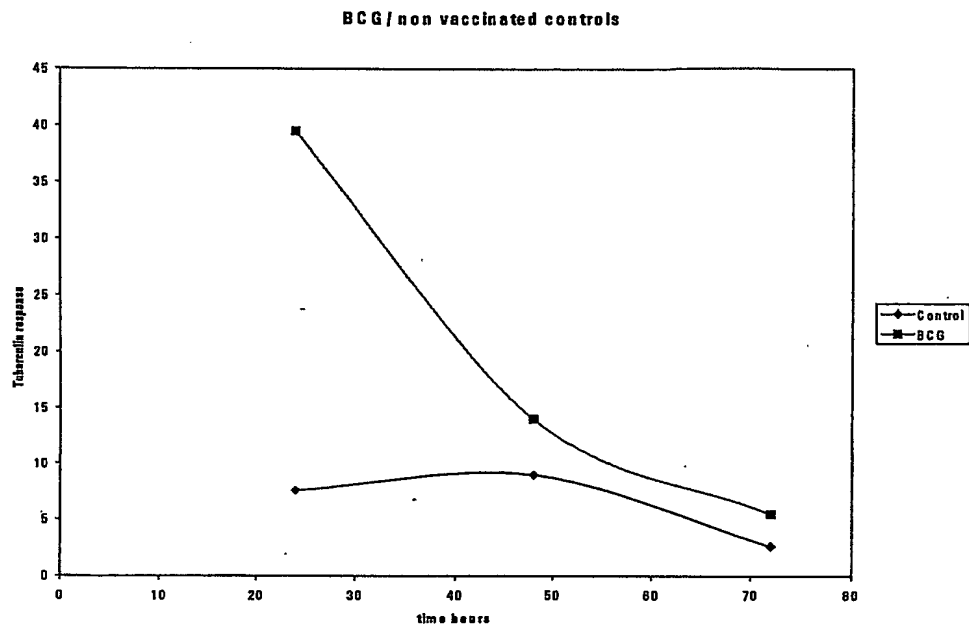


Figure 7

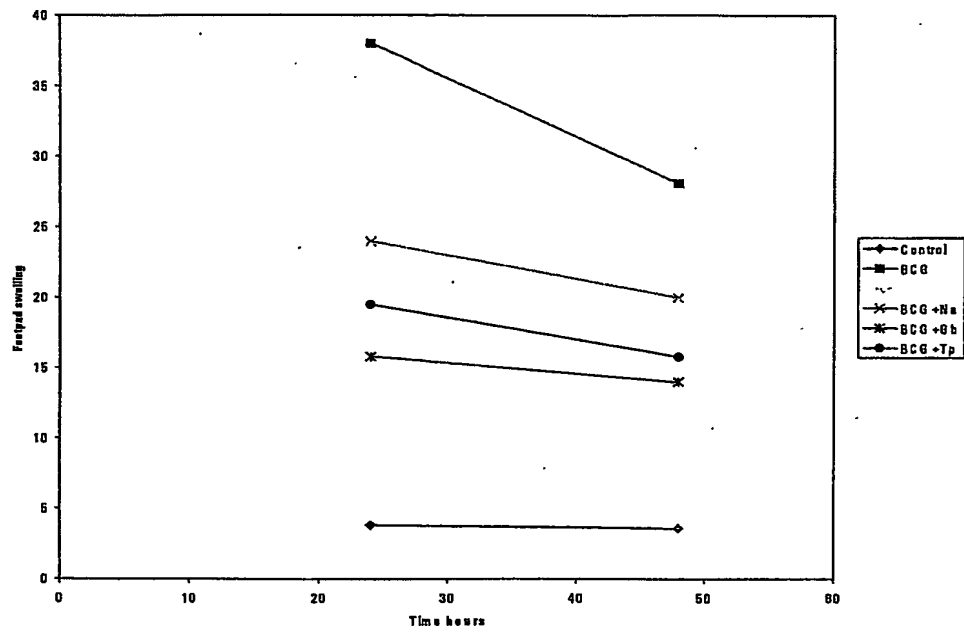


Figure 8

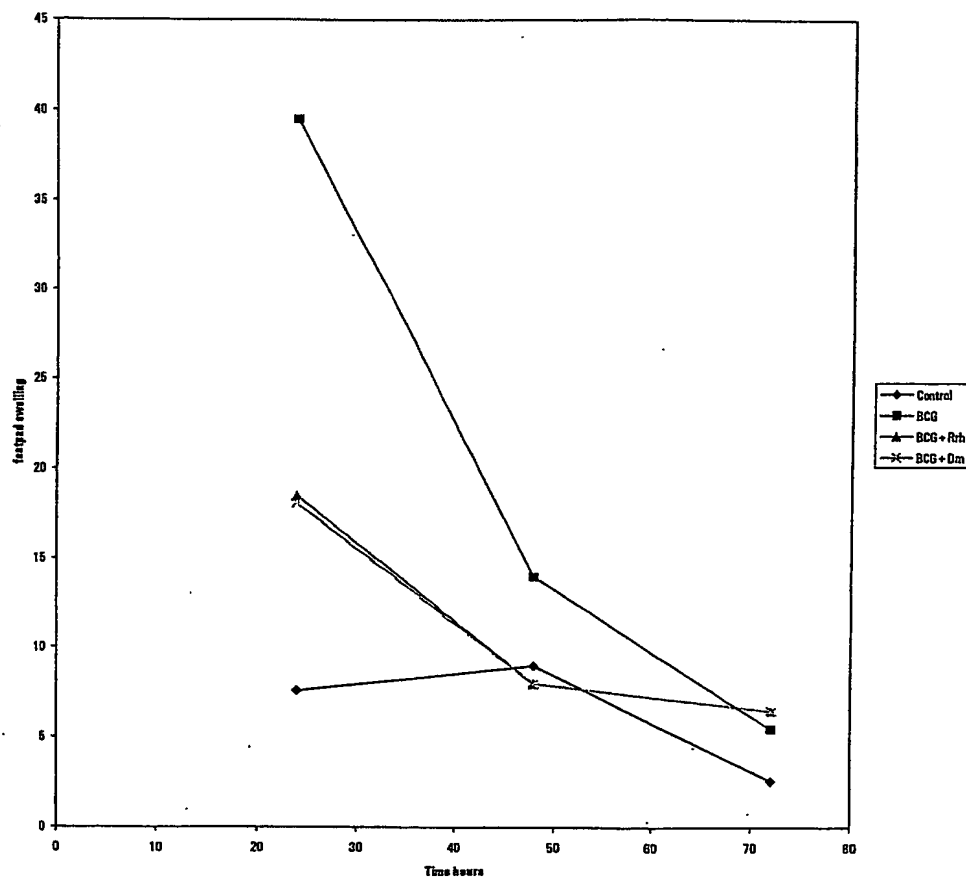


Figure 9

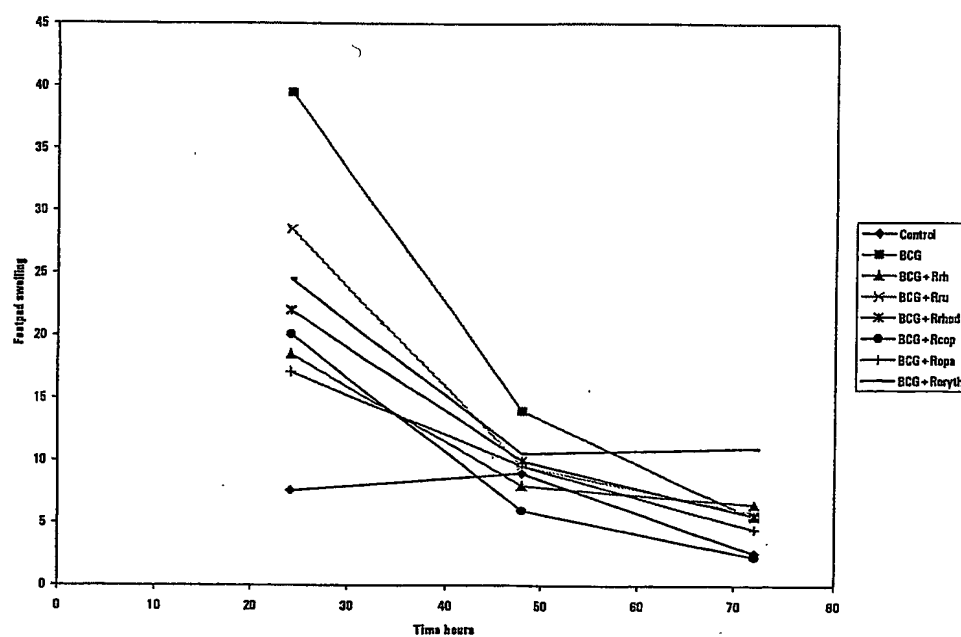
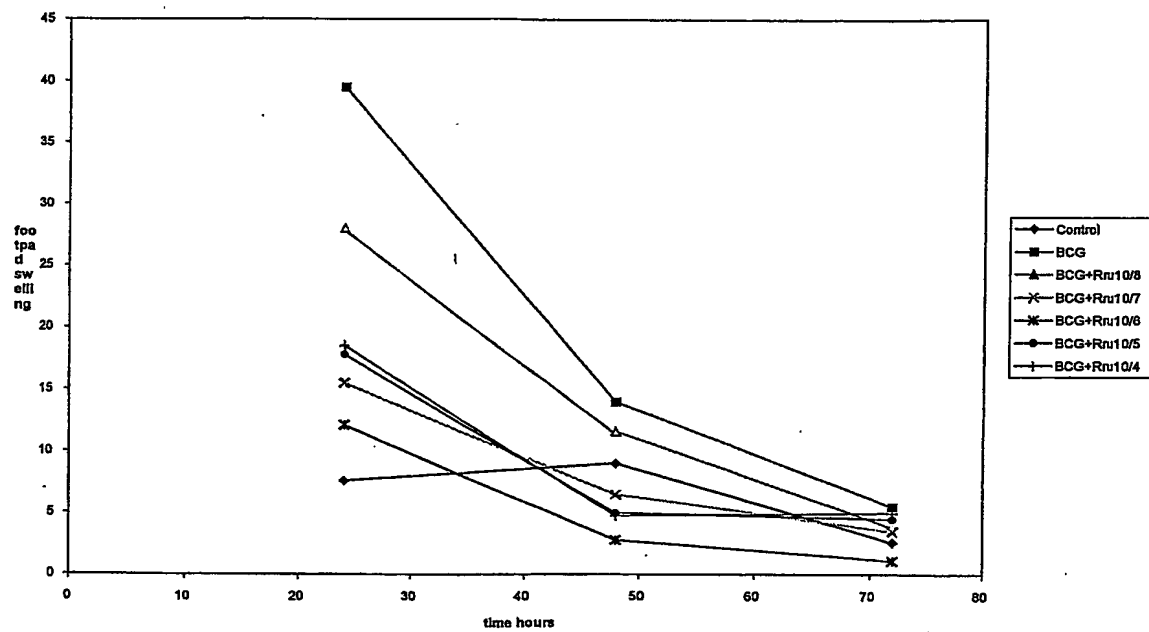


Figure 10

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